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Harrington Station Unit 2 Fabric Filter System
Special Testing Report

Southwestern Public Service Co.
Amarillo, TX

Prepared for

Industrial Environmental Research Lab.
Research Triangle Park, NC

Oct 81



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by

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ABSTRACT

A special test program was conducted by Southwestern Public Service Company to determine mass emissions of particulate, sulfur dioxides, and oxides of nitrogen from a large utility baghouse controlling emissions from a boiler firing low sulfur Western coal. The sampling was conducted in four phases. The first phase conducted in December 1978 verified that NO_x emissions, as measured by EPA Method 7, were not significantly affected by the baghouse. Emissions of SO_2 measured consistently below the New Source Performance Standard of $1.2 \text{ lb}/10^6 \text{ Btu}$ at a level of approximately $0.8 \text{ lb}/10^6 \text{ Btu}$. Particulate tests indicated an average emission rate of $0.028 \text{ lb}/10^6 \text{ Btu}$. Overall results of the special testing program indicate that Southwestern's Harrington Station, using low sulfur and low ash Western coal, can consistently operate within all applicable mass emission rates while using a fabric filter for air quality control.

CONTENTS

	<u>Page</u>
Abstract	
List of Figures	vi
List of Tables	viii
Conversion Table	ix
I. Introduction and Background	1
II. Conclusions	3
III. Harrington Station Description	4
IV. Equipment, Methods, Calibration Procedures	7
V. SPS Test Phases and Results	88
Appendices	
A. Coal Analysis	
B. Raw Field Data	
C. Field Notes	

LIST OF FIGURES

<u>Number</u>		<u>Page</u>
Figure IV-1	Lear Siegler Console	8
Figure IV-2	Pump Flow Diagram	9
Figure IV-3	Manual Sampling Probe	10
Figure IV-4	Probe Liner Supports	11
Figure IV-5	Method of Wrapping Nichrome Wire	12
Figure IV-6	Attachment of Pitot Tube Lines and Thermocouple Casing	13
Figure IV-7	Type S Pitot Tubes	14
Figure IV-8	Umbilical Cord	15
Figure IV-9	Probe Extension	17
Figure IV-10	Vertical Sampling Set-up	18
Figure IV-11	Stack Sampling Set-up	19
Figure IV-12	Form for Calibration of Standard Volume Flasks	20
Figure IV-13	Flask and Wet Test Meter Hook-up	22
Figure IV-14	Form for Wet Test Meter Calibration	23
Figure IV-15	Sampling Console and Wet Test Meter Hook-up	24
Figure IV-16	Form for Pre-Calibration Leak Check	26
Figure IV-17	Dry Gas Meter Calibration Form	27
Figure IV-18	Orifice Calibration Form	28
Figure IV-19	Form for Standardization of Thermometers	30
Figure IV-20	Magnehelic Connection to Inclined Manometer	31
Figure IV-21	Form for Magnehelic Calibration	33
Figure IV-22	Probe Heater Check Form	34
Figure IV-23	Pitot Tube Calibration Form	35
Figure IV-24	Orsat Bag Evacuation	37
Figure IV-25	Combustion Gas Sampling Train	39
Figure IV-26	Gas Analysis Data Sheet	40
Figure IV-27	Orsat Analyzer	41
Figure IV-28	Particulate Sampling Train	43
Figure IV-29	Particulate Data Sheet	44
Figure IV-30	Particulate Field Data Form	46
Figure IV-31	Moisture Fraction Form	47
Figure IV-32	Checklist for Recovery of Particulate Samples	48
Figure IV-33	Particulate Blank Results	50
Figure IV-34	Particulate Results	51
Figure IV-35	Acetone Purity Check Form	52
Figure IV-36	SO ₂ Sampling Train	55
Figure IV-37	SO ₂ Sample Field Data Form	57
Figure IV-38	Bottle Label	58
Figure IV-39	Checklist for Recovery of SO ₂ Samples	59
Figure IV-40	SO ₂ Analysis Form	60
Figure IV-41	SO ₂ Data Summary	61
Figure IV-42	Isooproponal Check for Peroxide Impurity	64
Figure IV-43	Standardization of Barium Perchlorate	65
Figure IV-44	Standardization of NaOH	66

Figure IV-45	Standardization of H_2SO_4	67
Figure IV-46	SO_2 Quality Assurance Samples	72
Figure IV-47	Assembly of NO_x Sample Train	74
Figure IV-48	Flask Stopcock Positions	74
Figure IV-49	NO_x Sampling Data Form	75
Figure IV-50	NO_x Analysis Sheet	77
Figure IV-51	Analysis of Standards for NO_x	81
Figure IV-52	Method 5 Dry Gas Meter Inter- Laboratory Survey	87
Figure V-1	Fabric Installation, July 1979	91

LIST OF TABLES

<u>Number</u>		<u>Page</u>
Table III-I	Fabric Filter System Design Parameters	5
Table IV-I	Dry Gas Volume Computer Program	69
Table IV-II	Program For Method 7 Calibrations	78
Table IV-III	Suggested Performance Criteria for Particulate Sampling Equipment	83
Table IV-IV	Suggested Performance Criteria for SO ₂ Sampling Equipment	84
Table IV-V	Suggested Performance Criteria for NO _x Sampling Equipment	85
Table V-I	Phase I Test Results	89
Table V-II	Phase II Test Results	92
Table V-III	Phase III Test Results	94
Table V-IV	Phase IV Test Results	95
Table V-V	Results of SPS/GCA Joint Test	97

CONVERSION TABLE

To Convert From	To	Multiply by
BTU/lb	joules/kg	4186.8
°F	°C	(°F-32)/1.8
°F	°C	(°F-32).56
ft	m	0.305
gr/scf	gm/m ³	2.29
in.	cm	2.54
in w.g.	mm Hg	1.87
lb	gm	454
miles	km	1.609
oz.	grams	28.350
ppm	mg/liter	1.0
Btu	joules	1055
cfm	cubic meters/hr	1.699
ft/sec	centimeters/sec	30.48
sq. ft.	sq. meters	0.093
lbs. of tension	newtons	4.448
stack pressure in Hg	kg/cm ²	0.035
wscf/10 ⁶ Btu	wscm/joule	2.6x10 ⁻¹¹
lbs/10 ⁶	nanogram/joule	433.3
lbs/wscf	micrograms/m ³	16.018x10 ⁸
ft ²	meters ²	0.093

I. INTRODUCTION AND BACKGROUND

Electric utilities, in recent years, have begun a search for the most efficient, economical method of emission control to use in association with boilers designed to burn low sulfur coal. One promising application is fabric filtering. Fabric filters have been used successfully by other industries for particulate capture. Southwestern Public Service Company (SPS), an electric utility based in Amarillo, Texas, first began considering baghouses for utility application in the mid-1970's after having installed a precipitator/wet scrubber on the company's first coal-fired unit. SPS compared costs, operating and maintenance parameters, and visited the only two utility baghouses in operation at that time (Nucla Station and Sunbury Station). Based on this limited information, SPS selected a fabric filter system to be used for emission control on its second coal-fired unit.

The SPS baghouse was the first to be installed on a new utility boiler firing low sulfur Western coal; therefore, its operation and performance was of special interest to the U. S. Environmental Protection Agency (EPA). In 1977, the EPA executed a contract with SPS to conduct an in-depth study of this new emission control alternative. The objective of the five-year program is to fully characterize the performance of a large utility baghouse controlling emissions from a boiler firing low sulfur Western coal.

One of the major aims of the SPS/EPA study is to determine mass emissions of particulate, sulfur dioxides, and oxides of nitrogen. The contract specifies a special testing program designed to measure these parameters and relate results to the collection efficiency of the fabric filter system.

The original work plan called for three special tests by SPS. However, the test dates were altered in order to allow for changes in operating conditions and budget limitations. Tests were conducted by SPS at the following times:

Phase I	December 11-15, 1978
Phase II	October 8-12, 1979
Phase III	March 24-25, 1980
Phase IV	August 26-29, 1980

The contract also called for three more specialized tests to be conducted by an outside consultant under subcontract to Southwestern. GCA Corporation was selected to sample simultaneously at five locations for the following parameters: particulate, $C_7 - C_{17}$ organic compounds, $C_1 - C_6$ organic compounds, CO_2 , O_2 , CO , SO_2 , SO_3 , NO_x , and particle size distribution. GCA conducted their first test in February 1979. Results from this test led SPS and the EPA to a decision to reduce this phase of the test program to only one more test to be performed as a joint test by SPS and GCA. Furthermore, the test parameters were reduced to the following: particle size distribution by three methods, and SO_2 and SO_3 sampling at three locations. This series of tests was performed in June, 1980. Results of the GCA tests will be reported in a separate report.

Even though it was the intent of SPS to accomplish the objectives of the tasks set forth in the contract, several problems were experienced during initial operation of the baghouse which kept it from being classified as a typical or standard type of air quality control device.

These problems centered around the control of the cleaning cycle, control of the deflation pressure during the cleaning cycle, proper tension on the bags when they were first installed, a higher than normal bag failure rate, and indications that Harrington Station's fly ash had some unique characteristics which made predicting proper design difficult. Programs were developed to investigate these problems and to attempt to find solutions.

II. CONCLUSIONS

The sampling conducted by Southwestern Public Service Company on Harrington Station Unit 2 was conducted in four phases. The first phase conducted in December 1978, verified that NO_x emissions, as measured by EPA Method 7, were not significantly affected by the fabric filter system. The unit operated below the applicable New Source Performance Standard (NSPS) of $0.7 \text{ lb}/10^6 \text{ Btu}$ at a level of approximately $0.64 \text{ lb}/10^6 \text{ Btu}$. The concentrations of NO_x were tested simultaneously at two inlet, two outlet and one stack location. Since the NO_x emissions were determined to be less than the NSPS and stack emissions were close to the inlet emissions, it was concluded that no significant removal efficiencies were occurring within the fabric filter system. Therefore, the sampling for oxides of nitrogen was discontinued after the first test.

Sulfur dioxide emissions were determined by EPA Method 6 four times during the test program. Stack emissions of SO_2 agreed very well with the theoretical SO_2 emission. The unit operated consistently below the NSPS of $1.2 \text{ lb}/10^6 \text{ Btu}$ at a level of approximately $0.8 \text{ lb}/10^6 \text{ Btu}$. Based upon the observed inlet and stack concentration there was no appreciable absorption of SO_2 on the filter cake during filtering.

Particulate emissions were actually determined by EPA Method 5 five times during the test program. The unit was tested during October 1979, with fabric that was approximately 3 months old. This test gave the lowest average emission of $0.020 \text{ lb}/10^6 \text{ Btu}$. Tests conducted later averaged an emission rate of $0.028 \text{ lb}/10^6 \text{ Btu}$ with extreme emission rates of 0.016 to $0.044 \text{ lb}/10^6 \text{ Btu}$.

Tests conducted during the fabric filter study indicate that Harrington Station, using low sulfur and low ash Western coal, can consistently operate within all applicable mass emission rates while using a fabric filter system for air quality control.

III. HARRINGTON STATION DESCRIPTION

Harrington Station, located approximately five miles northeast of Amarillo, Texas, is owned and operated by SPS. The facility consists of three 360 MW pulverized, coal-fired boilers. Unit 1 is equipped with an electrostatic precipitator followed by a marble bed wet scrubber. Units 2 and 3 are each equipped with a fabric filter system.

The Unit 2 steam turbine is a Westinghouse designed, single reheat, tandem compound, double flow turbine with 28.5-inch last stage blades. The steam turbine drives a 400 MVA electric generator.

The steam boiler is a Combustion Engineering steam generator, tangentially fired, capable of producing 2,682,383 pounds of steam per hour at 2500 psig, 1005° F superheat, and 1005° F reheat. The basic fuel source for Harrington is pulverized Western coal, but the steam generator furnace is also designed for alternate fuels (natural gas and No. 2 burner fuel oil).

Coal is delivered to Harrington by rail from an Atlantic Richfield Company mine in Campbell County, Wyoming. Average parameters of the coal are 8,475 Btu/lb, 0.3% sulfur, and 5.5% ash. (A representative coal analysis for each test phase is shown in Appendix A.) Coal is fed with automatic equipment into coal mills where it is crushed to approximately 200 mesh. This fine coal dust is blown into the boiler furnace with high temperature air where combustion takes place.

The emission control device selected for Unit 2 is a Wheelabrator-Frye, Inc. fabric filter baghouse system. The baghouse is designed to operate at a flue gas flow of 1,650,000 acfm at 313° F. Minimum design efficiency is 98.6%, which would permit 0.1 lbs. of particulate/million Btu out the stack. The exterior of the baghouse has 3.5 inches of fiberglass insulation; there is no insulation between plenums and compartments.

The baghouse is equipped with bypass dampers for start-up, emergency operation, and shut-down. Other design parameters are summarized in Table III-1.

There are two baghouses on Harrington Unit 2, one designated east and one designated west. Each one has its own operating control system and all bypass dampers are separated for each system.

Bag cleaning is accomplished by a deflate/shake cycle. During the normal filtering sequence the outlet and the reinflation dampers are open, while the deflation damper is closed. During the cleaning cycle the reinflation damper and outlet damper is closed, leaving the inlet damper open. After an initial settle period, the deflation damper opens, which provides just enough reverse air to gently collapse the bags, causing the filter cake to fracture. The cake is then separated from the fabric by mechanical shaking. After a final settle period the reinflation and outlet dampers open, putting the compartment back in service.

The Unit 2 baghouse has four cleaning cycle modes which were designed to provide maximum flexibility of operation. Mode 1 will clean the west baghouse and then the control system will reset; Mode 2 will clean the east baghouse and reset; Mode 3 will clean compartments 1 through 28 before the system is reset; and Mode 4 will clean the east and west baghouses simultaneously, one compartment at a time on each baghouse.

Table III-1. Fabric Filter System Design Parameters

Compartments	28
Bags per Compartment	204
Model	No. 366, Series 11.5 RS Dustube
Bag Diameter	11.5 inches
Bag Length	30 feet, 8 inches
Bag Spacing, Center to Center	14.0 inches
Design Air-to-Cloth Ratio, Gross	3.16:1
with 1 compartment down	3.27:1
with 2 compartments down	3.40:1
Bag Reach	2
Cleaning Mode	Deflate/Shake
Total Cloth Area	5.14 x 10 ⁵ ft ²
Bag Design	Caps with eye bolts
Original Fabric	Criswell fiberglass, silicone/graphite coating

SPS brought the Harrington Station Unit 2 baghouse into service on June 21, 1978. Approximately three weeks after the baghouse was initially started, SPS was able to operate Unit 2 at full load with only coal in service. The unit has operated at loads consistently above 200 Mw and during peak periods, it has handled loads of 367 Mw.

Since start-up, a variety of adjustments have been made, including a complete change-out of the bags. Shaker mechanisms have been re-worked and timing sequences have been altered. These changes took place at different periods in the special testing program. They will be discussed in more detail and their relationship to a particular phase of testing more specifically defined in Section V of this report. For a more detailed overview of the Harrington Station baghouse performance, refer to the first and second annual reports (1, 2).

IV. EQUIPMENT, METHODS, CALIBRATION PROCEDURES

A. Introduction

The procedures described in this section were implemented by the Environmental Group of the System Laboratory for collection, recovery, and analysis of samples for the determination of particulate, SO_2 , NO_x , and combustion gases (O_2 and CO_2) in flue gases during special test periods. Also discussed are methods for fabrication of equipment, calibration procedures, and the quality control/quality assurance procedures employed by SPS for the special testing program. Samples of raw field data are included in Appendix B.

This section of the special testing report will be somewhat familiar to those experienced in stack sampling. The procedures are basically the same as EPA Methods 3, 5, 6, and 7 as outlined in the Federal Register.

B. Fabrication of Equipment

1. Sampling Consoles: The SPS sampling team utilized five sampling consoles during the test program. Three of the consoles were purchased from Lear Siegler and the general specifications of these consoles are given in Figure IV-1. The two remaining consoles were fabricated by System Lab personnel and are similar in design to standard EPA approved consoles except that the vacuum pumps are located externally. The pumps are plumbed according to the drawing in Figure IV-2.

2. Probes and Probe Liners: The System Lab has six probes with steel liners. Four of these are for baghouse testing and have 316 stainless steel liners. The liners are 1/2-inch outside diameter (O.D.). Two of the probes are 22 feet long (inlet) and two are 32 feet long (outlet). The other two probes are for stack testing. They have 5/8-inch O.D. Inconel liners. The probe used for the special testing program is 12 feet long. Drawings showing the assembly of the probes are included here in Figures IV-3 and IV-4.

The probe liners have an 18/9 male ball joint on one end. The liners are wrapped twice with fiberglass tape, then electrically insulated 25 gauge nichrome wire is spiraled the length of the probe and two more layers of fiberglass tape wrapped over that. On the stack probes, one 50-foot piece of nichrome wire is used. The two 22-foot probes have two 50-foot sections of nichrome wire connected in parallel. The two 32-foot probes have two 25-foot sections and one 50-foot section of wire connected in parallel. Figure IV-5 shows the method of wrapping the nichrome wire. A ground wire is attached to each probe liner to prevent electrical shock in case of a short. The hook-up wire used is 16 gauge, Teflon-insulated hook-up wire. The pitot tube lines and thermocouple casing (copper tubing) are attached to the probes in the manner shown in Figure IV-6.

3. Type S Pitot Tubes: The type S pitot tubes used for special testing are constructed out of 1/4-inch O.D. stainless steel according to the specifications given in the Federal Register of August 4, 1978. They are approximately 18 inches long and have Swagelok fittings for attachment to the pitot tube lines. A drawing of these tubes can be found in Figure IV-7.

4. Umbilical Cord: Figure IV-8 is an end view of the umbilical cord used on the stack. The pitot tube lines are 1/4-inch inside diam-

SAMPLING CONTROL UNIT **LEAR SIEGLER CONSOLE**

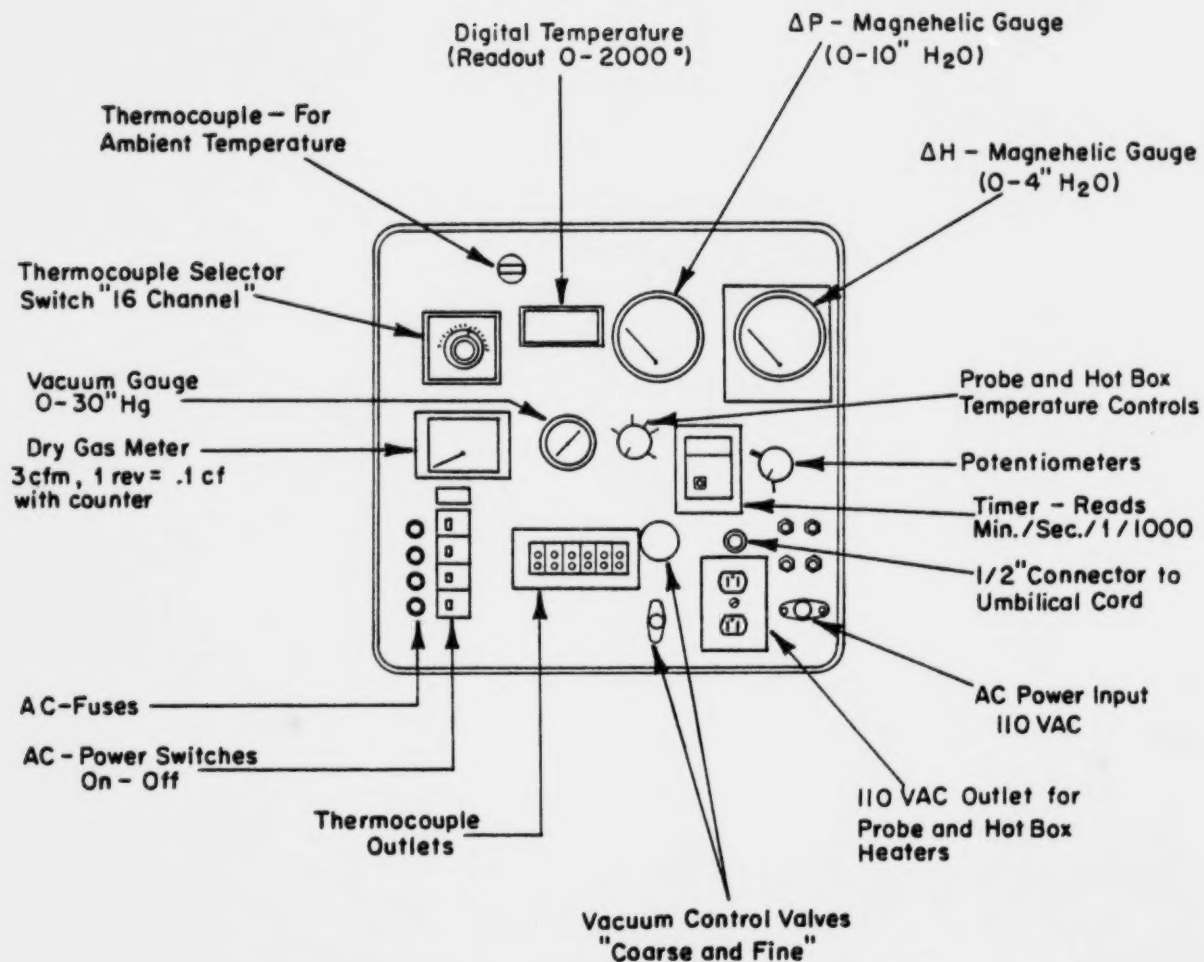


FIGURE IV-1. LEAR SIEGLER CONSOLE

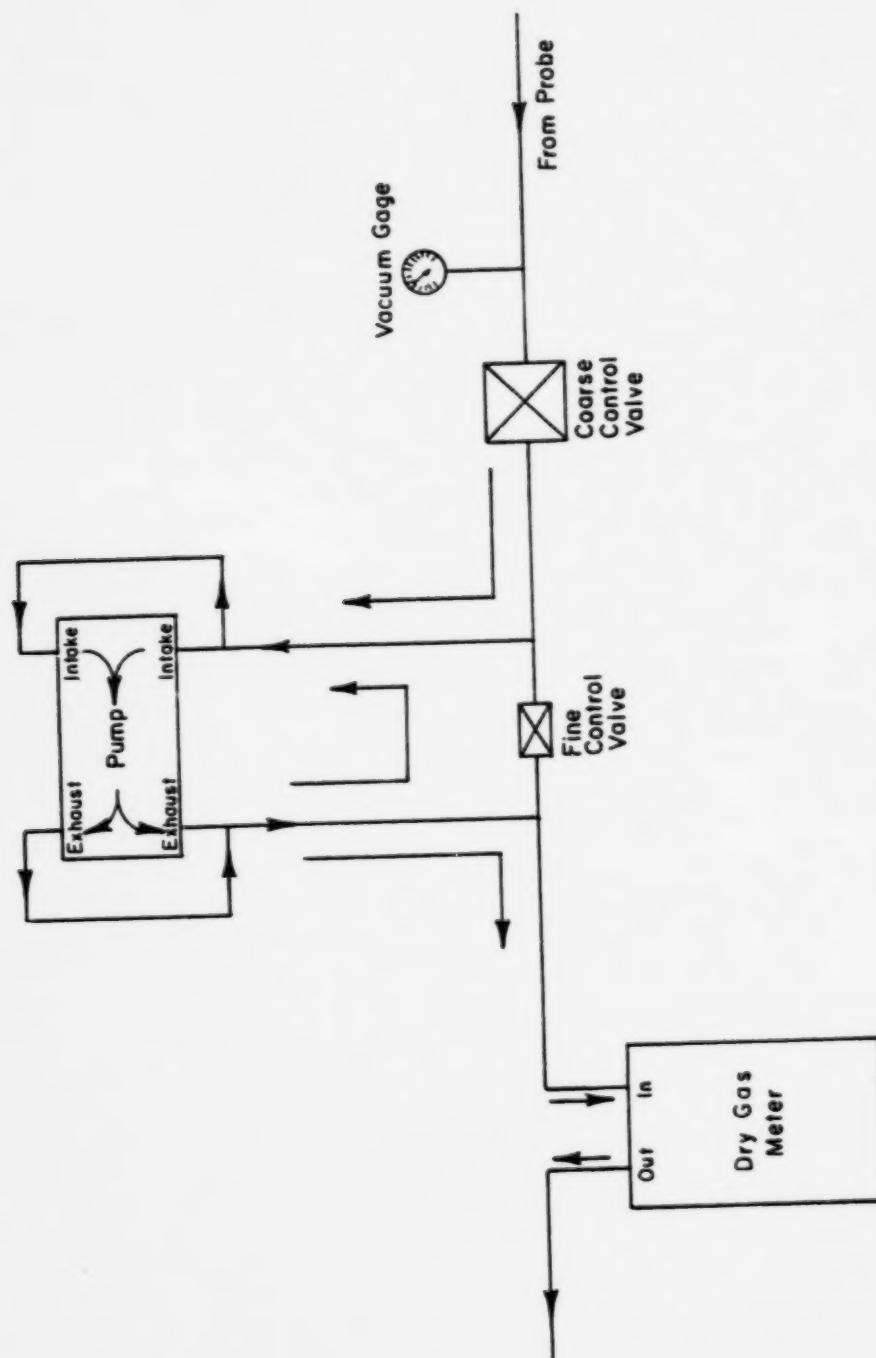
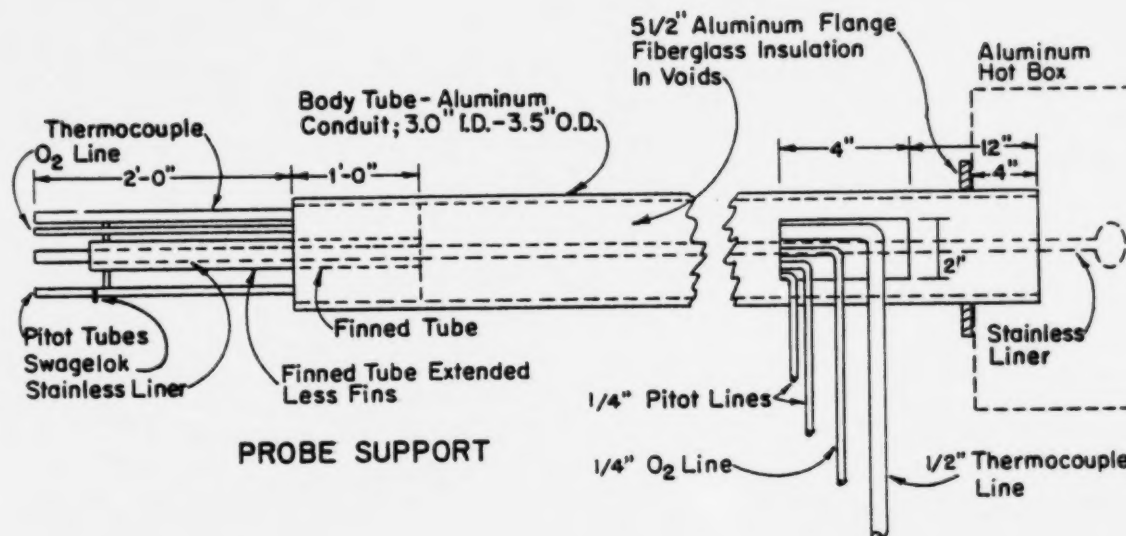
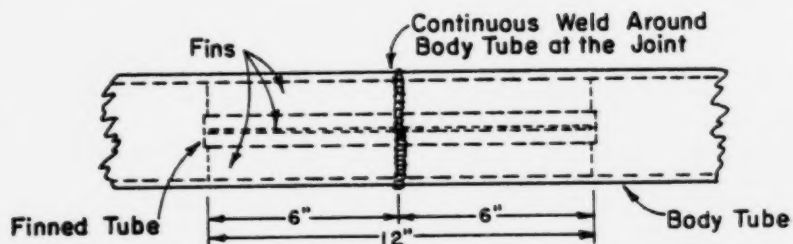


FIGURE IV-2. PUMP FLOW DIAGRAM



2 - 22 Foot Probes
2 - 32 Foot Probes



TYPICAL JOINT DETAIL

Note: Fins continuously welded to finned tube, spot welded to body tube at end of fins.

FIGURE IV-3. MANUAL SAMPLING PROBE

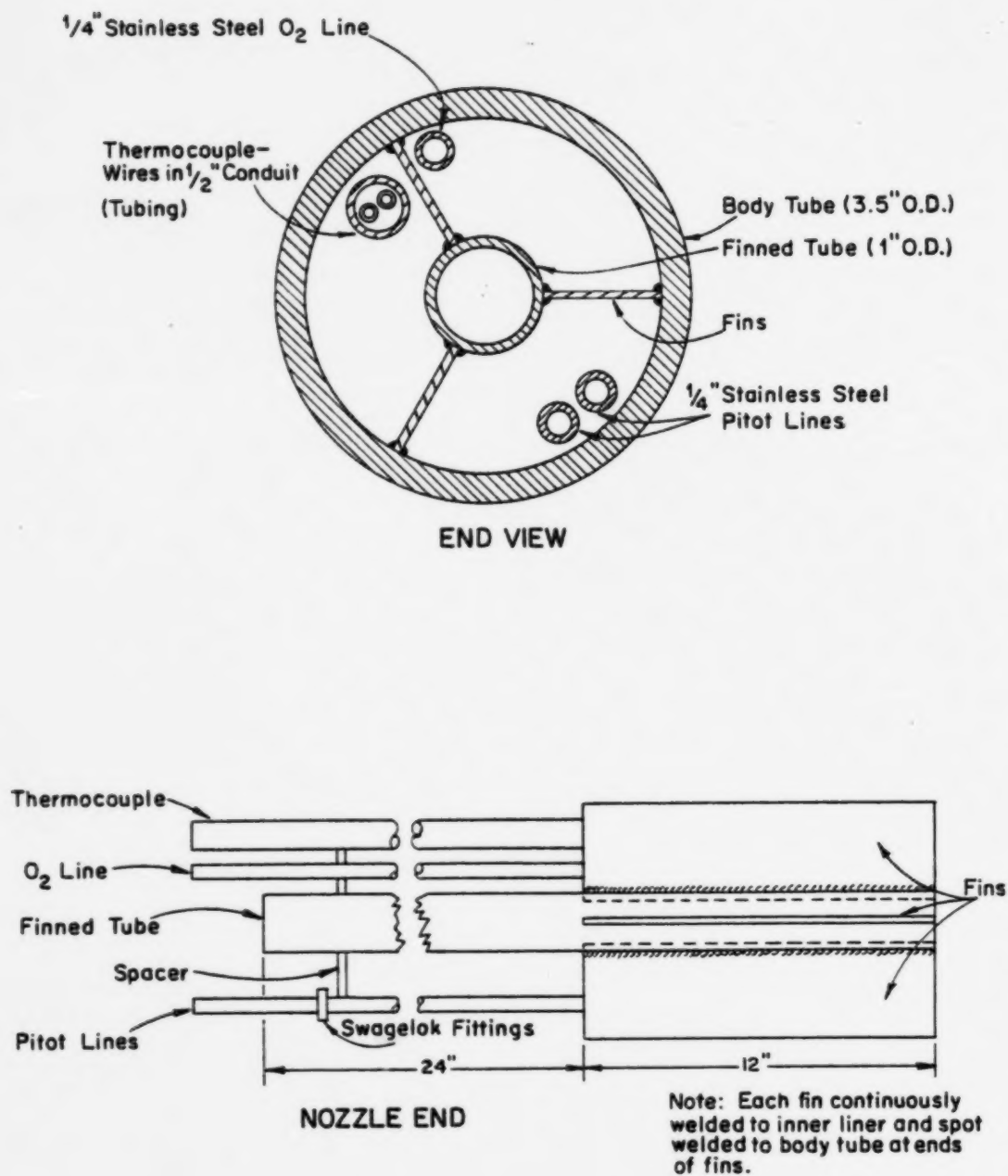


FIGURE IV-4. PROBE LINER SUPPORTS

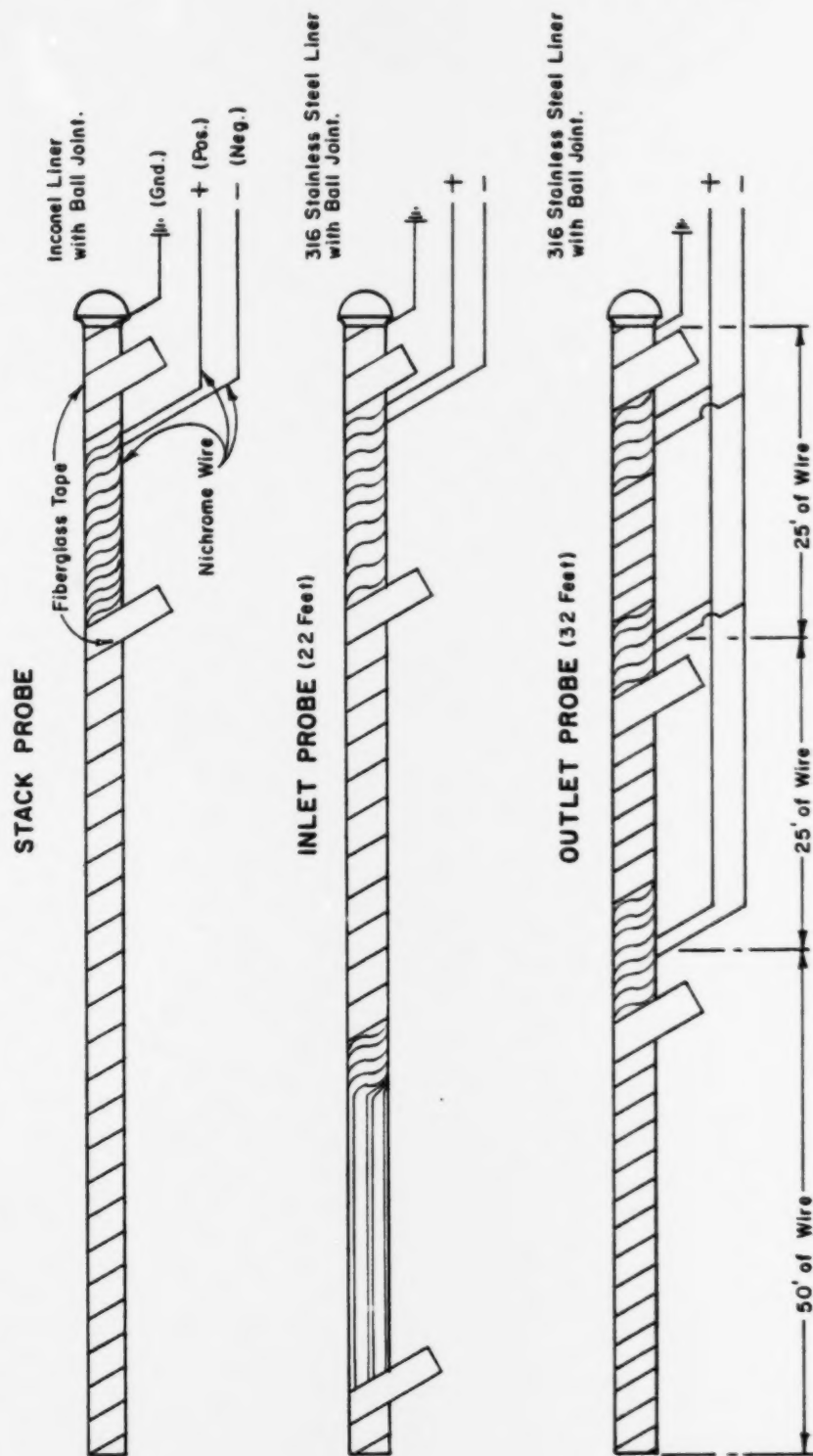


FIGURE IV-5. METHOD OF WRAPPING NICHROME WIRE

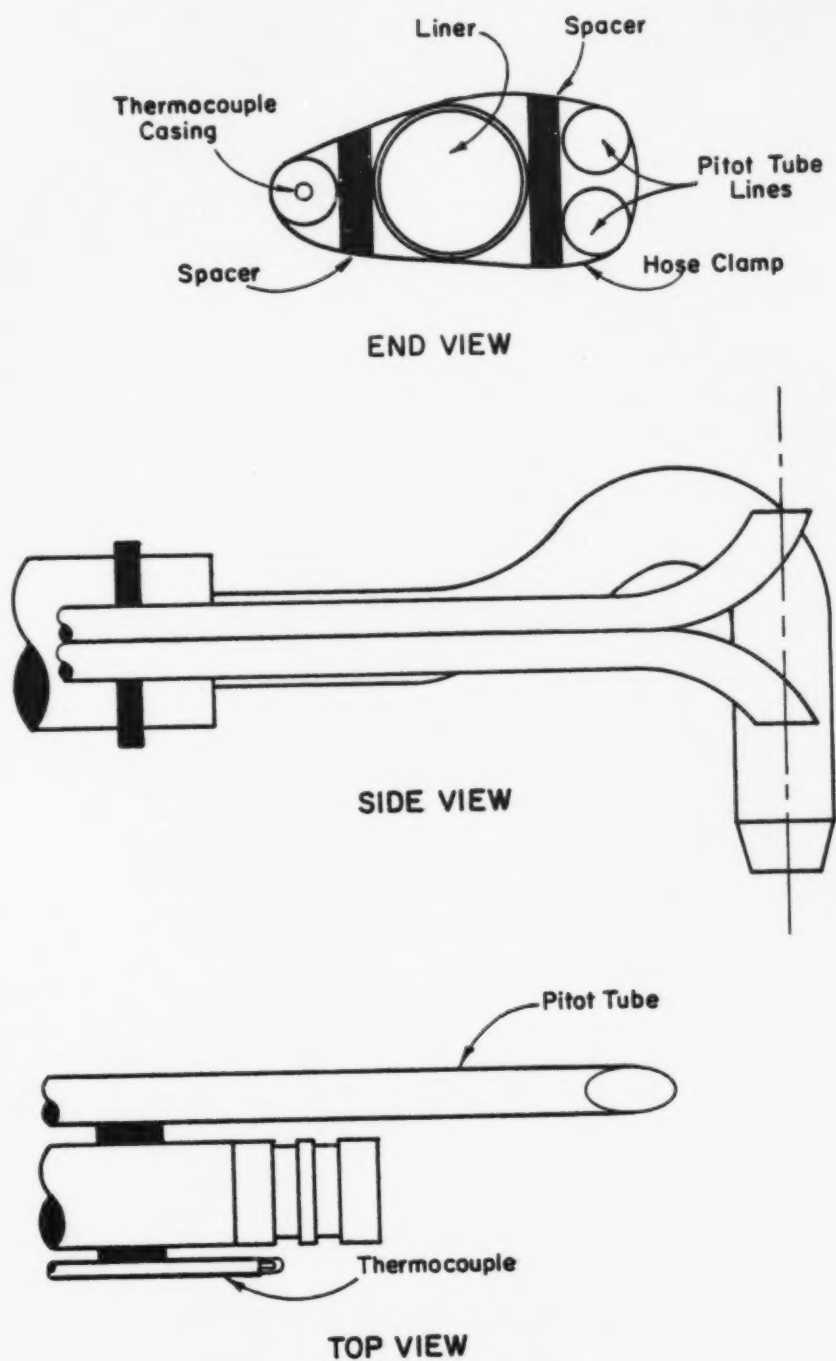
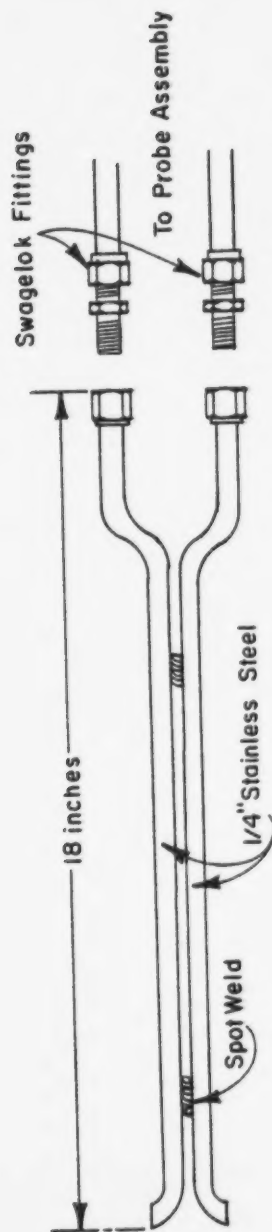


FIGURE IV-6. ATTACHMENT OF PITOT TUBE LINES AND THERMOCOUPLE CASING



Note: Pitot tube lines and probe lines are color coded to maintain correct orientation - red and green.

FIGURE IV-7. TYPE S PITOT TUBES

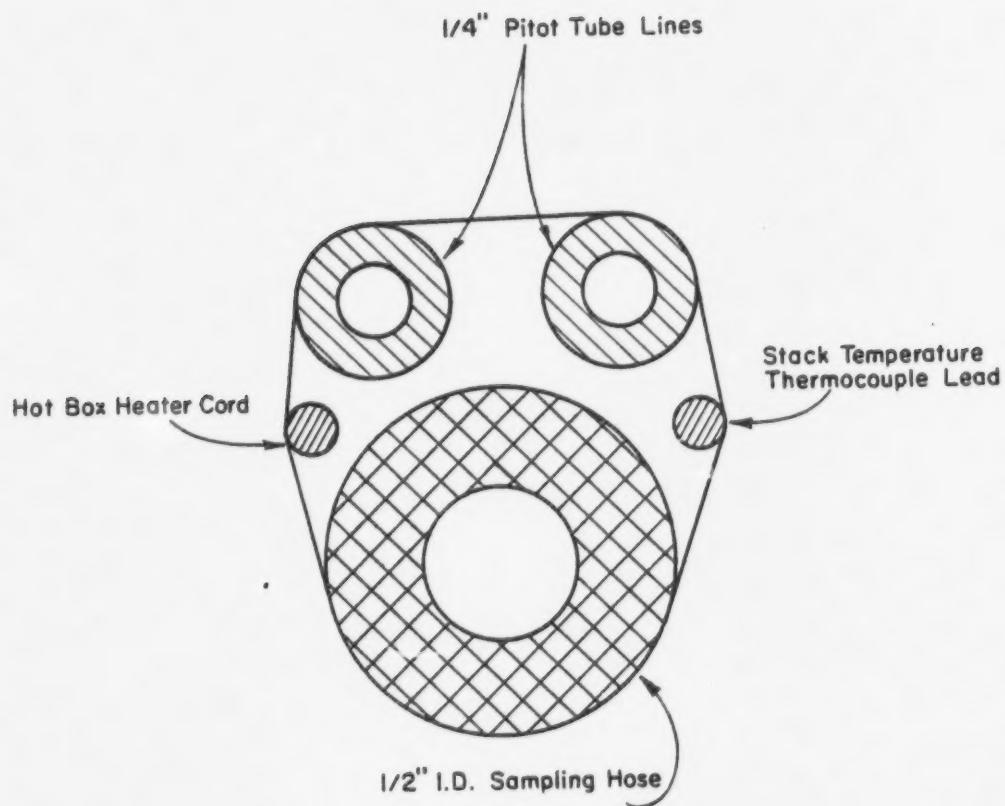


FIGURE IV-8. UMBILICAL CORD

eter (I.D.) welding tubing. The sampling hose is 1/2-inch I.D. thick-wall Tygon tubing. These are taped together at about 1-foot intervals with black electrical tape. The pitot tube lines have 1/4-inch O.D. stainless steel tubing inserted in the end that goes to the sampling console and Tygon tubing attached to make connection to the console magnehelics easier. All tubing to hose connections are made fast with hose clamps.

5. Probe Extensions: The extensions of the probe from the hot box to the cold box for vertical sampling is Teflon, high-temperature heated hose. The hose has a built-in electric heating element that can provide uniform heat up to 450° F. throughout the entire length of the hose. The core is made of 1/2-inch I.D. Teflon. The hose has half an inch of thermal insulation. A drawing of this probe extension can be found in Figure IV-9. A drawing of a complete vertical sampling set-up and a complete stack sampling set-up can be found in Figures V-10 and IV-11.

C. Calibration Procedures

1. Calibration of Standard NO_x Volume Flask: The standard volume flasks were calibrated for special testing by filling the flask to the mark with D.I. water and then weighing the water to the nearest gram. The temperature of the water and the density of water at that temperature was also recorded. (See Figure IV-12, Form for Calibration of Standard Volume Flasks.) This procedure was repeated three times. The ml of water used was determined by the following formula:

$$(xg) \frac{(1)}{e} = ml \quad (1)$$

where

xg = weight of water used to the nearest gram

e = density of water at a given temperature.

This volume was then converted to cubic feet using the following formula:

$$ml \times \frac{ft^3}{28,316 \text{ ml}} = ft^3 \quad (2)$$

where

ml = ml of water as determined above

28,316 = #ml/ft³.

The three values were averaged and that value assigned to the flask as the standard volume when filled to the mark.

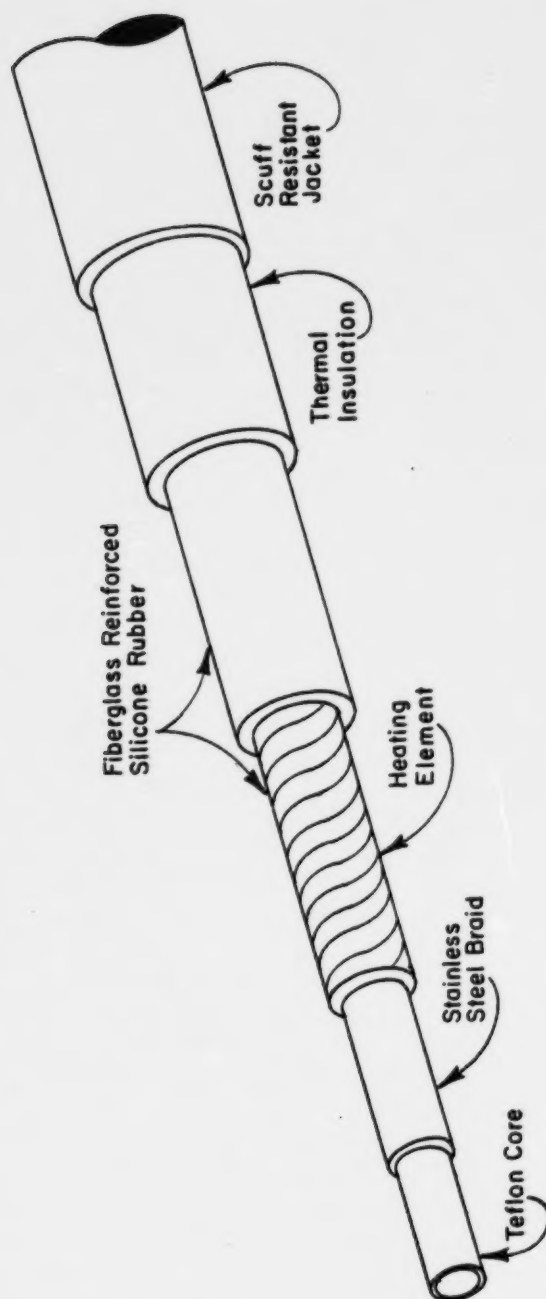


FIGURE IV-9. PROBE EXTENSION

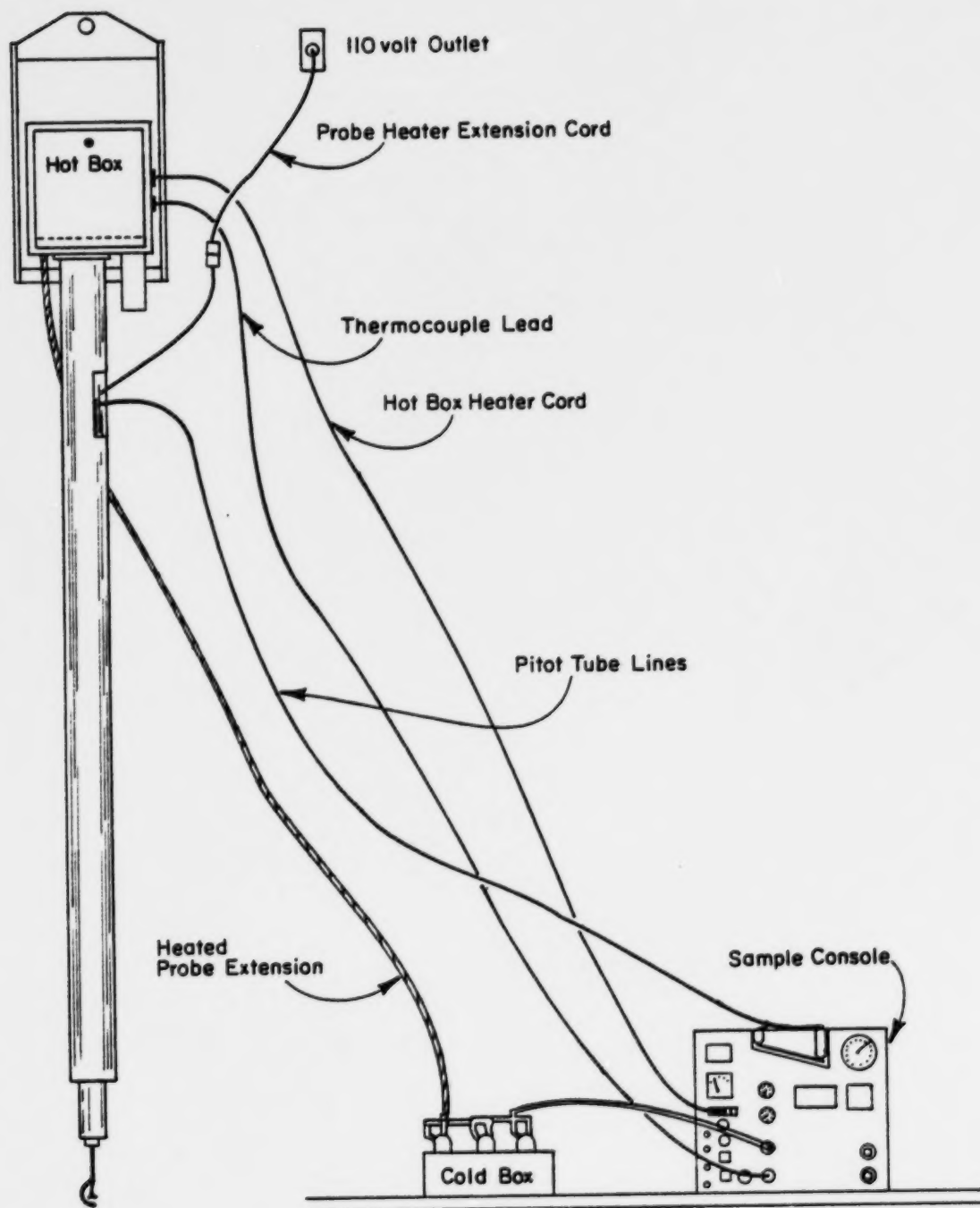


FIGURE IV-10. VERTICAL SAMPLING SET-UP

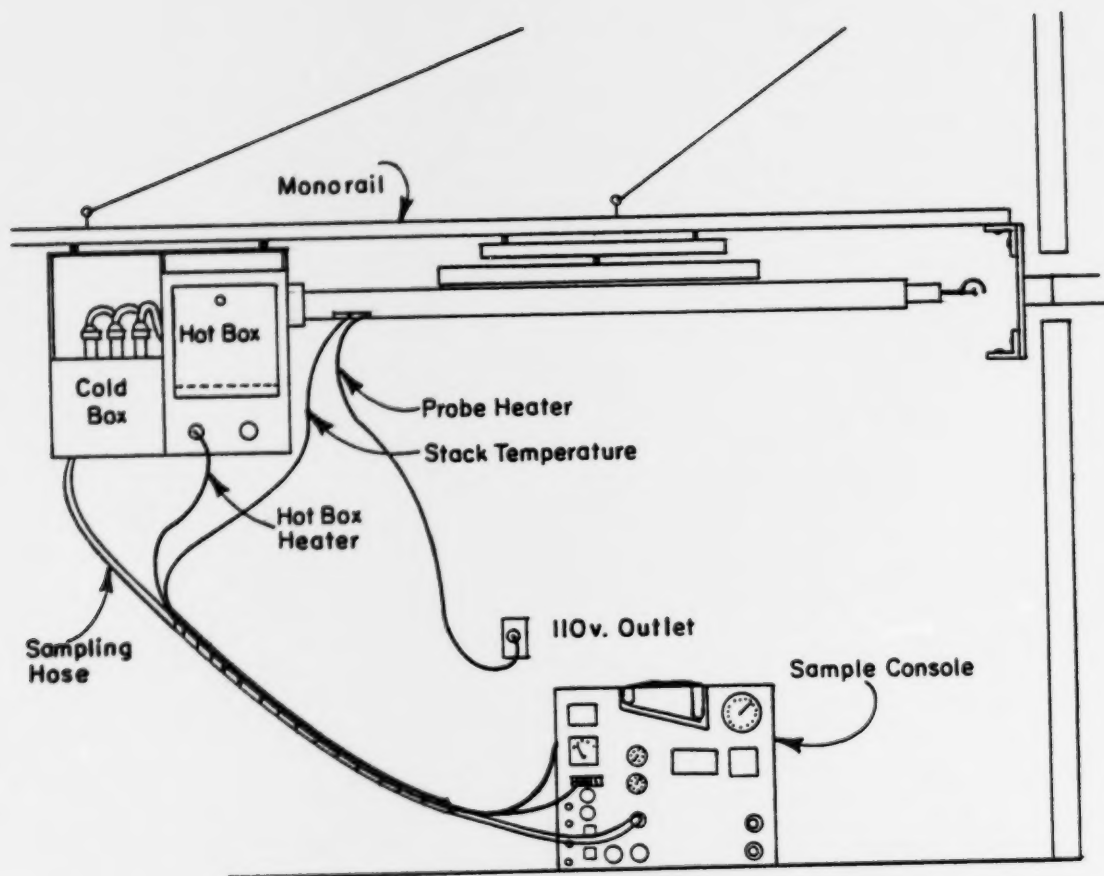


FIGURE IV-11. STACK SAMPLING SET-UP

Calibration of Standard Volume Flasks

Flask # _____

Date _____

	Weight of Water	Temperature	Density (e) at Given Temperature
Run #1			
Run #2			
Run #3			

$$(xg)\left(\frac{1}{e}\right) = \text{ml}$$

$$\text{ml} \times \frac{\text{ft}^3}{28,316 \text{ ml}} = \text{ft}^3$$

Standard ft³

Run #1

Run #2

Run #3 _____

Avg.

Figure IV-12. Form for Calibration of
Standard Volume Flasks

2. Calibration of Wet Test Meter: The wet test meters were calibrated during the test program by passing a known volume of saturated gas through them. To do this, air was displaced by a known volume of water in a leak-free system through the wet test meter.

A 46-liter flask was used to displace the air into the wet test meter. The exact volume of air that was displaced was determined by calibration of the flask. The flask and wet test meter were hooked up as shown in Figure IV-13. The water level in the wet test meter was correct and the initial wet test meter reading recorded. (See Figure IV-14, Form for Wet Test Meter Calibration.) D.I. water was used to displace the air. The water was added to the flask at a fairly constant rate. The level of water was stopped exactly on the mark that is shown on the flask. After filling the flask, the final wet test meter reading was recorded. The difference in the final and initial readings was calculated. The wet test meter correction factor was calculated using the following formula:

$$\text{Wet Test Meter Correction Factor} = \frac{\text{standard volume}}{\text{volume measured by wet test meter}} \quad (3)$$

The value obtained and the date was put on a label and placed on the wet test meter.

3. Calibration of Dry Gas Meters: The calibration of the dry gas meter (DGM) was done by making simultaneous total volume measurements with a calibrated wet test meter and the dry gas meter. The complete sampling console system was leak checked before the calibration was done. For metering systems with diaphragm pumps, the normal leak-check procedure will not detect leakages within the pump.

To do the pre-calibration leak check, the sampling console and wet test meter were hooked up as shown in Figure IV-15. The wet test meter (WTM) was leveled and the outlet of the WTM connected to the inlet line of the console. The outlet of the bubbler was connected to the WTM inlet. A 10-minute calibration run was done at a rate of 0.02 cfm. The initial wet test meter and dry gas meter readings were taken as were the final readings at the end of the run. The following formula was used to calculate the leak rate:

$$\frac{\text{Vol. Measured by Wet Test Meter} - \text{Vol. Measured by Dry Gas Meter}}{10} \quad (4)$$

This value must be less than 0.02 cfm.

After a good pre-calibration leak check was obtained, the calibration of the dry gas meter was continued. Set-up for calibration was the same as for the leak check procedure. The sampling pump was turned on and allowed to warm up for a few minutes. The pump was valved off and the initial DGM and WTM readings were taken. The pump was then valved on and the rate adjusted to about 0.2 cfm and the vacuum to about 10 inches of Hg. The initial ambient inlet to DGM and outlet to DGM temperatures were also recorded.

The pump was allowed to run until 10 (or more) cubic feet had passed through the dry gas meter. Temperature readings were taken right before stopping the pump, then the pump was stopped and meter readings were recorded. The calibration factor was determined by using the following formula:

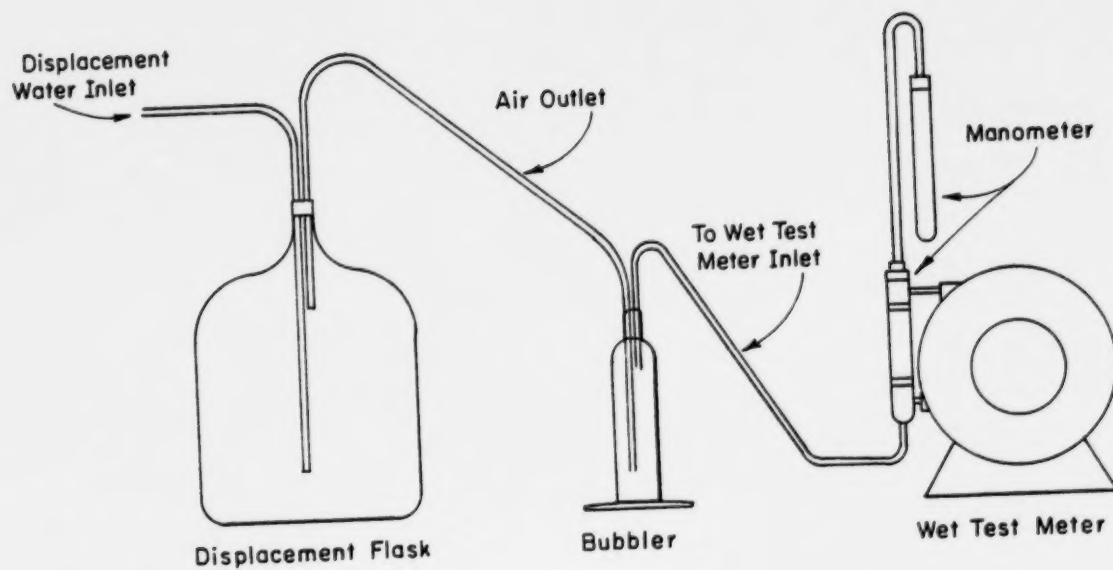


FIGURE IV-13. FLASK AND WET TEST METER HOOK-UP

Wet Test Meter Calibration

Wet Test Meter No. _____

Date _____

Run No. _____

Calibrated by _____

Standard Flask No. _____

Standard ft.³ _____

Wet Test Meter Reading Final _____

Wet Test Meter Reading Start _____

Wet Test Meter Reading Difference (WTMRD) _____ ft.³

Wet Test Correction Factor = $\frac{\text{Standard}}{\text{WTMRD}}$ = _____

Figure IV-14. Form for Wet Test Meter Calibration

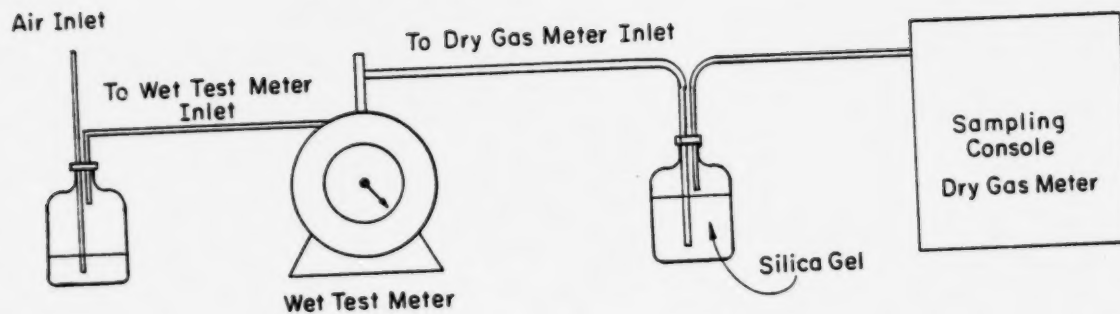


FIGURE IV-15. SAMPLING CONSOLE AND WET TEST METER HOOK-UP

$$\text{DGM} = \frac{(\text{WTD}) (P \text{ Bar} \times \frac{\Delta H}{13.6}) (\text{WF}) (T_m + 460)}{(\text{DFG} - \text{DGI}) \text{DGI} (P \text{ Bar} + 13.6 \frac{\Delta P}{Ta + 460})} \quad (5)$$

where

WTD = wet test meter difference

WF = wet test meter correction factor

DGF = final DGM reading

DGI = initial DGM reading

Ta = average wet test meter temperature

Tm = average dry gas meter temperature

ΔH = inches of water across DGM orifice

ΔP = inches of water across wet test meter.

This procedure was repeated three times and the average of the three runs assigned as the correction factor of the DGM. The date of calibration and the correction factor was put on a label and placed on the sampling console. (See Figure IV-16, Form for Pre-Calibration Leak Check and Figure IV-17, Dry Gas Meter Calibration Form.)

4. Orifice Calibration: The orifice was calibrated by passing various known flow rates through it and recording the pressure differential between upstream and downstream taps on the Orifice Calibration Form (Figure IV-18).

The procedure for the actual calibration was as follows: The barometric pressure and the dry gas meter correction factor was recorded on the data sheet. The pump was turned on and the vacuum adjusted to 10 inches Hg and the delta H to 0.2 inches of H₂O. Initial entrance and exit temperatures of the DGM were recorded. The length of time in seconds required for one cubic foot of air to pass through the DGM was measured. This number and the final entrance and exit temperature were recorded. The above procedure was repeated for each 0.2 increment of delta H up to 3.0 inches, and then for every 0.5 increment.

The flow rate for each increment in standard cubic feet per minute (SCFM) was calculated using the following equations:

$$\text{Vol in ACFM} = \frac{\text{DG}_f - \text{DG}_i}{\text{time}} \times 60 \quad (6)$$

$$\text{Vol in SCFM} = (\text{Vol in ACFM}) (\text{DGMCF}) \frac{(530)}{T+460} \frac{(\text{Bar. Pressure})}{29.92} \quad (7)$$

Pre-Calibration Leak Check

Console# _____

Date _____

Checked by _____

Final Wet Test Meter Reading _____

Initial Wet Test Meter Reading _____

Wet Test Meter Difference (WTMD) _____

Final Dry Gas Meter Reading _____

Initial Dry Gas Meter Reading _____

Dry Gas Meter Difference (DGMD) _____

WTMD - DGMD must be 0.02 DVM

10

Figure IV-16. Form for Pre-Calibration Leak Check

Dry Gas Meter Calibration Form III

By Wet Test Meter

Run # _____

Date _____

$\Delta P =$ _____ IN. H_2O

Control Terminal _____

$\Delta H =$ _____ IN. H_2O

Purpose _____

$P_{bar} =$ _____ IN. Hg.

Calibrated _____

Ambient _____ °F

Wet Test Meter, Final (WTMF) _____

_____ cf

Wet Test Meter, Initial (WTMI) _____

_____ cf

WTD = (WTMF-WTMI) _____

_____ cf

Calibration Factor For Wet Test Meter (WF) _____

Date of Last Calibration For Wet Test Meter _____

Temperatures

Wet Test Meter

Dry Gas Meter

Inlet

Outlet

Initial

Final

Sum

2 ÷

_____ °F

+

Average WTM

Temp(T_a)

_____ °R

4 ÷

_____ °F

_____ °F

Average Dry Gas Meter Temp (T_m) _____

_____ °R

Dry Gas Reading, Final (DGF) _____

_____ cf

Dry Gas Reading, Initial (DGI) _____

_____ cf

(DGF-DGI) = DF

_____ cf

Dry Gas Meter Factor (DF) = _____

WTD(P + $\frac{\Delta H}{13.6}$)(WF) (T_m)

DF(P + 13.6) (T_a) =

() (+ $\frac{13.6}{13.6}$) (x) = _____

() (+ $\frac{13.6}{13.6}$) ()

Figure IV-17. Dry Gas Meter Calibration Forms

[illegible]

$$\text{Vol in ACFM} = \frac{DG_f - DG_i}{\text{Time}} \times 60$$

$$\text{Vol in SCFM} = (\text{Vol in ACFM}) \left(\frac{530}{\frac{T}{1} + 460} \right) \left(\frac{\text{Bar. pressure}}{29.92} \right)$$

FIGURE IV-18. ORIFICE CALIBRATION FORM

where

ACFM = actual cubic feet per minute
SCFM = standard cubic feet per minute
DG_f = final dry gas meter cubic feet
DG_i = initial dry gas meter cubic feet
DGMCF = dry gas meter correction factor
T = average meter temperature.

5. Standardization of Thermometers: Dial-type thermometers are calibrated using in-glass mercury thermometers that had been certified. At least three points were used to check the thermometers. (These are the freezing and boiling points of water and room temperature.) The standard thermometer number, standard thermometer reading, and each thermometer reading was recorded on the data sheet. (Figure IV-19, Form for Standardization of Thermometers.) The dial type thermometers must agree to within ± 1.5 percent of the standard value.

6. Calibration of Magnehelic Gauges: The procedure for calibrating magnehelic gauges involved comparison of value for pressure differentials on the magnehelic and an inclined manometer. The magnehelic is tied in parallel with an inclined manometer as shown in Figure IV-20. The impact side of the manometer was attached to the intake side of the orifice and the static side to the exhaust side of the orifice. The pump was started with the coarse valve open and the manometer disconnected. The coarse valve was then closed and the manometer connected. The valve was then opened slowly and the delta H as measured by the manometer was compared with that of the magnehelic. The flow was adjusted to give delta H readings in 0.5 inch increments through the range of the magnehelic.

Standardization of Thermometers

Date: _____

Standard #	Standard Reading	_____	_____	_____	_____
------------	------------------	-------	-------	-------	-------

Figure IV-19. Form for Standardization of Thermometers

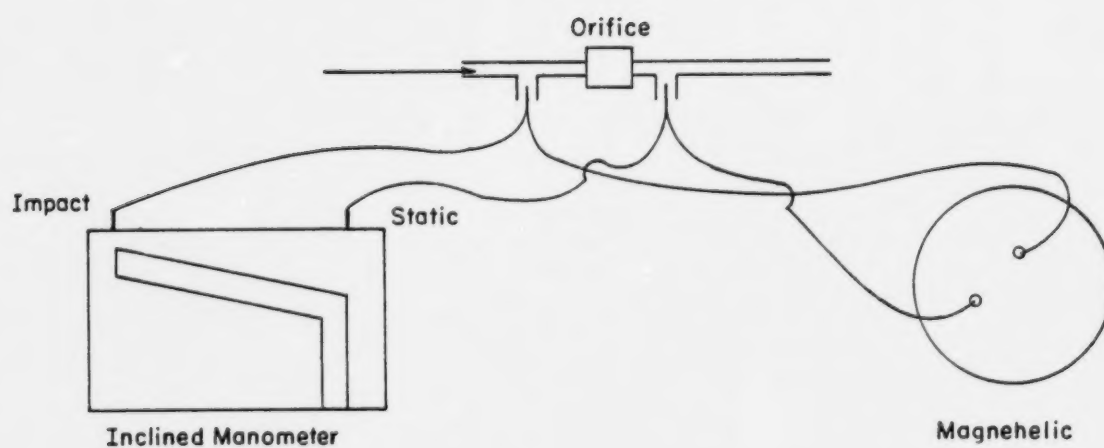


FIGURE IV-20. MAGNEHELIC CONNECTION TO INCLINED MANOMETER

If delta P magnehelics were being calibrated, tubes were attached to the suction or pressure of the magnehelic and either blown or sucked on the line and clamped off. The magnehelic must agree to within at least 5 percent of what the manometer reads. Figure IV-21 is the form for recording magnehelic calibrations.

7. Probe Heater Checks: The probe heaters were checked by measuring the internal temperature of the probe liner as a function of time. To do this, a piece of thermocouple wire long enough to reach about the middle of the probe was crimped at the ends and inserted in the liner. The temperature was recorded every minute or so until the rate of increase slowed. Then it was recorded about every 10 minutes until the maximum temperature was reached. (See Figure IV-22, Probe Heater Check Form.)

8. Calibration of Pitot Tubes: If pitot tubes could not be calibrated in a wind tunnel, the following procedure was used for calibration. The stack on Nichols Station Unit 3, located one mile from Harrington Station, was used to provide the flow for calibration. The type S pitot tubes were calibrated against a standard pitot tube. Two inclined manometers were used for the calibration. The manometers were leveled and zeroed. Each pitot tube was then hooked up to a manometer and both systems were leak checked. After good leak checks were obtained, the two tubes were inserted into the stack and aligned in the same plane. The delta P, as measured by each pitot tube, was recorded (Figure IV-23, Pitot Tube Calibration Form). The two tubes were then moved to another position in the stack and the readings taken again. This procedure was conducted at least three times. The type S pitot tube was then rotated and the procedure repeated for the other side of the tube. The pitot tube coefficient was calculated using the formula below:

$$C_p = C_p(\text{std}) \frac{p_{\text{std}}}{p} \quad (8)$$

where

C_p = Type S pitot tube coefficient

C_p = Standard pitot tube coefficient (use 0.99 if not known)

p_{std} = velocity head measured by the standard pitot tube

p = velocity head measured by the type S pitot tube.

This calculation was done for each of the three runs on each side of the pitot tube. The average of the three runs is the pitot tube coefficient assigned to the tube for that side.

Magnehelic Calibration

Console # _____

Date _____

Calibration done by _____

Manometer Reading

Magnehelic Reading

Figure IV-21. Form for Magnehelic Calibration

Probe Heater Check

Date: _____

Probe # _____

Time

Probe Internal Temperature

[illegible][illegible]

Checked by: _____

Figure IV-22. Probe Heater Check Form

PITOT TUBE CALIBRATION

(using a wind tunnel)

PITOT TUBE NUMBER _____ DATE _____

CALIBRATED BY _____

STANDARD PITOT TUBE NUMBER _____

STANDARD PITOT TUBE CALIBRATION FACTOR _____

WIND TUNNEL FLOW	STANDARD PITOT TUBE TUBE ΔP in inches H ₂ O	"S" TYPE PITOT TUBE in inches H ₂ O	C _p "S" TYPE CALIBRATION FACTOR
0.3" H ₂ O			
0.4			
0.5			
0.6			
0.7			
0.8			
0.9			
1.0			
1.5			
2.0			

0.3" H₂O

0.4

0.5

0.6

0.7

0.8

0.9

1.0

1.5

2.0

$$C_p = C_{pstd} \left(\frac{\Delta P \text{ STANDARD}}{\Delta P \text{ "S" TYPE}} \right)^{1/2}$$

$$C_p \text{ avg.} = \frac{\sum C_p}{n}$$

$$C_p \text{ avg.} = \frac{\quad}{10}$$

$$C_p \text{ avg.} = \frac{\quad}{\quad}$$

C = "S" Type Calibration Factor.

C_{pstd}. = Standard Pitot Tube Calibration Factor.

C_{pavg}. = Average "S" Type Calibration Factors.

FIGURE IV-23. PITOT TUBE CALIBRATION FORM

SPS Environmental Date 5-19-80

For the coefficient factor to be acceptable, the average deviation of the three run values must be 0.01 and the absolute value of the difference of the two sides must be 0.01. The calculation of the average deviation is as follows:

$$\text{average deviation} = \frac{\sum_{i=1}^3 C_p - \bar{C}_p}{3} \quad (9)$$

where

C_p = the type S tube coefficient for a given run and given side

$\bar{C}_p$ = the average tube coefficient.

D. Collection and Analysis For Combustion Gas

The gas analysis apparatus used in the special test program for gas analysis, (i.e., O_2 and CO_2), is a Burrell standard Orsat analyzer. The Orsat bags used to collect the gas sample are 5-liter, "snout" type, constructed of layers of polyester, polyvinyl chloride, aluminum foil, polyimide, and polyethylene. The probe used to sample is 1/4-inch stainless steel and the pump used is a one-way squeeze bulb. Reagents used are DiSorbent (CO_2 absorption) and Oxsorbent (O_2 absorption).

1. Pre-sampling Preparation: Prior to sampling, equipment was inspected, cleaned, and checked for proper operation. Orsat bags to be used for sample collection were evacuated (see Figure IV-24) in the following manner:

Flexible tubing was connected to hose bib on the bag and the valve was opened on the bag. Using a hand pump, the bag was evacuated completely. While tubing was still connected to the hose bib, the valve was shut off and the tubing disconnected.

Leak checks of sampling trains were performed at sample locations just prior to sampling in the following manner. With probe tip plugged (or capped), the sample train was evacuated until the squeeze bulb collapsed. If the squeeze bulb remained collapsed for at least 30 seconds, the sample train was considered "leak free" from probe tip to discharge side of the squeeze bulb. With an Orsat bag unattached to the sample train, the flexible tubing was clamped or "pinched off" where it attached to the Orsat bag hose bib; the bulb with net was inflated by using a squeeze bulb. If the bulb with net remained inflated for at least 30 seconds, then the sample train was considered "leak free" from the discharge side of squeeze bulb to the point where flexible tubing connected to the Orsat bag.

2. Sample Collection: Without disconnecting the sample train, the probe was placed in the stack, making sure the probe tip was at least 3.5 feet into the stack, (i.e., sample point should be at least 3.5 feet from inner wall of stack). The probe was secured to the port opening to keep it from falling or vibrating excessively. The port opening was

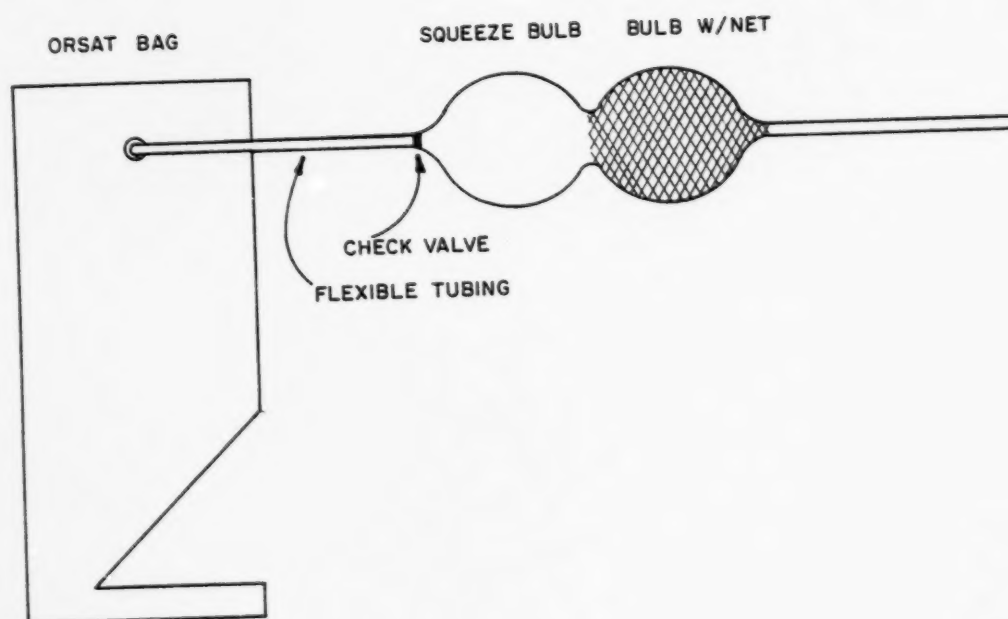


FIGURE IV-24. ORSAT BAG EVACUATION

covered with rags, duct tape, etc., to prevent ambient air from possibly diluting the sample. The sample train was then purged by using a squeeze bulb. After sufficient purging, (i.e., sample train completely clear of ambient air), pumping continued and the tip of the flexible tubing that connects to the Orsat bag was "pinched off". With the sample train completely assembled, (see Figure IV-25), the hose bib on the Orsat bag was opened and the bag was filled with sample gas. When the Orsat bag was full, the flexible tubing was disconnected. All pertinent data was recorded on the Gas Analysis Data Sheet (see Figure IV-26).

3. Sample Analysis: Prior to analyzing gas samples, a leak check was performed on the Orsat analyzers (see Figure IV-27) in the following manner. On each absorbing solution pipette, the absorbing solution liquid level was brought to the reference mark on capillary tubing by using a leveling bottle and then closing the pipette stopcock. Next, the leveling bottle was raised sufficiently to bring the liquid meniscus onto the graduated portion of the burette, closing the manifold stopcock. The meniscus position was recorded and observed; the level in each absorbing solution pipette was also observed over a 4-minute period (40 CFR Part 60, Appendix A, Method 3). If the liquid level in each absorbing solution pipette did not fall below the bottom of the capillary tubing and meniscus in the burette did not change by more than 0.2 ml during this 4-minute period, the Orsat analyzer was considered leak free.

Next, the liquid level in each absorbing solution pipette was determined to be at the reference mark. The manifold stopcock was set in position #2 and the water level in the burette was raised to the reference mark and the manifold stopcock was set in position #3. The hose bib on the Orsat bag was connected to the Tygon tubing on the Orsat manifold stopcock, and the hose bib on the Orsat bag was opened. The Tygon tubing was purged by applying pressure to the Orsat bag.

After purging, the manifold stopcock was set to position #4. By lowering the leveling bottle, the gas sample was drawn into a glass manifold and burette until meniscus was at the zero reference mark and then the manifold stopcock was set to position #1. The gas sample was run into the CO_2 absorbing solution pipette by opening the pipette stopcock and raising the leveling bottle (being careful not to run absorbing solution in the pipette past the reference mark). Next, the gas sample was run back into the burette.

Three complete passes were made. After the third pass, the absorbing solution level was run to the reference mark on the pipette and the pipette stopcock was closed. The leveling bottle water level was set at "even" with meniscus in the burette and the percent CO_2 was read directly from the graduated burette. A couple of more passes were made through the CO_2 absorbing solution and read again to ensure that CO_2 had been completely absorbed, i.e., until two consecutive readings were the same. The CO_2 reading was recorded.

The above procedure was repeated for O_2 . The reading from the burette included the percent CO_2 . The percent CO_2 was subtracted from this reading to get the percent O_2 , and then recorded. If more than ten passes were required to obtain a constant reading for any component gas, the absorbing solution in that respective pipette was changed out.

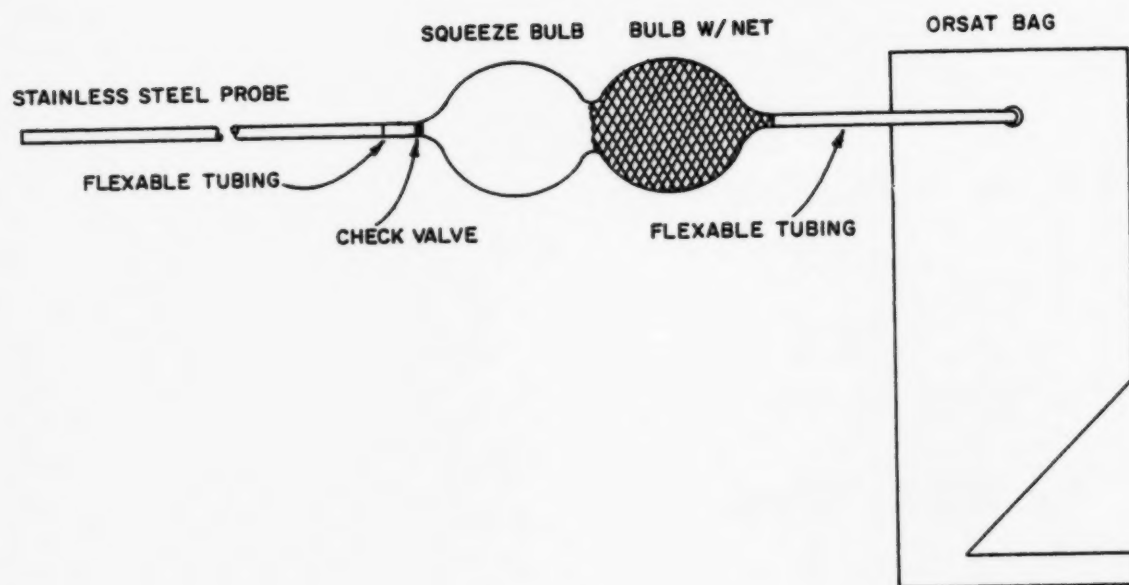


FIGURE IV-25. COMBUSTION GAS SAMPLING TRAIN

GAS ANALYSIS DATA

Plant/Location _____ Date _____
 Purpose _____ Sample No. _____

ORSAT ☐

FYRITE ☐

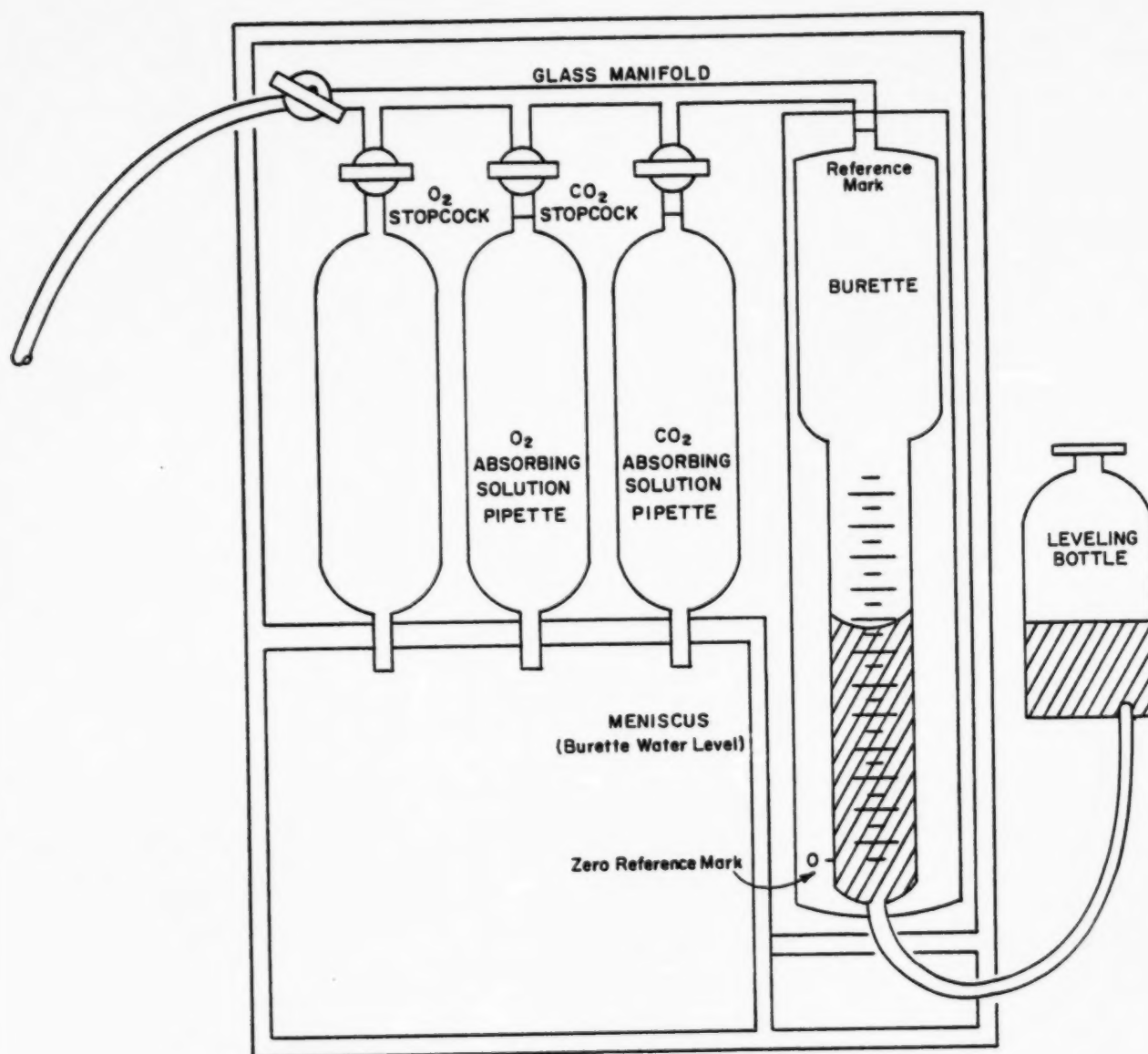
Bag No.	Time	Leak Ck.	CO ₂	O ₂	CO	
						Average
						Average
						Average

Sampled by _____

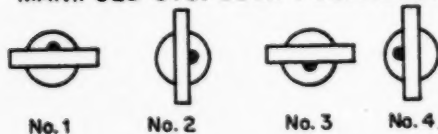
Analyzed by _____

FIGURE IV-26. GAS ANALYSIS DATA SHEET

ORSAT ANALYZER



MANIFOLD STOPCOCK POSITIONS:



- No. 1 - Closed.
- No. 2 - Vents glass manifold to atmosphere.
- No. 3 - Vents Tygon line to atmosphere.
- No. 4 - Vents glass manifold to Tygon line.

PIPETTE STOPCOCK POSITION

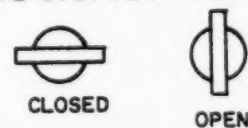


FIGURE IV-27. ORSAT ANALYZER

This complete analysis was repeated until three consecutive analyses varied by no more than 0.5 percent (absolute) by volume for each gas component being analyzed. The final results, i.e., average of the three consecutive runs, were reported on a dry volume percentage basis.

4. Quality Assurance: A calibration check was performed before every third field test or before any field test in which the Orsat apparatus had not been checked during the previous three months. To check the O_2 absorbing reagent and the operator technique, the percent O_2 in air was determined. The average of three replicates should be 20.8 ± 0.7 percent. A higher than average reading will normally indicate operator error and a lower than average reading will indicate leaks, exhausted O_2 absorbing reagent, and/or operator error.

Orsat bags were periodically checked for leaks in the following manner. The bag was inflated with ambient air by using a hand pump. The Orsat hose bib was connected to a water-filled U-tube manometer. The hose bib was opened and pressure reading (should be at least two inches H_2O) was recorded. It was allowed to stand for at least 10 minutes and if pressure remained constant, the Orsat bag was considered leak free. All calibration checks with dates, dates reagents were changed out, dates Orsat was used, and dates that Orsat bags were leak checked were recorded in the Orsat Log Book.

E. Collection and Analysis For Particulate

1. Sample Collection: The impinger train used for the special testing program was set up with four Greenburg-Smith impingers. Each impinger had a modified tip. The first and second impingers were to have known amounts of water, the third empty, and the fourth silica gel. The weights of all four impingers were determined to the nearest 0.5 grams before use. A collection bottle was filled with some of the D.I. water and labeled "D.I. Water Blank". All openings where contamination might occur were kept covered until just prior to being assembled for use.

Using tweezers and rubber gloves, a pre-weighed, labeled filter was placed in the filter holder with the numbered side down. The filter apparatus was assembled, making sure the gasket was in the proper position. The particulate sampling train was assembled as shown in Figure IV-28.

A preliminary and post leak check was performed in the following manner. After the sampling train was completely assembled or disassembled, the probe tip was plugged with a rubber stopper and the control valve to the pump opened. A vacuum of at least 15 inches of Hg for a preliminary leak check and a vacuum equal to or greater than the maximum vacuum reached during sampling for a post leak check was pulled. The needle on the dry gas meter was observed for any movement after the vacuum was reached. A leak rate not to exceed two percent of the average sampling rate was acceptable; however, if the rate was not zero, all connections were checked. The plug was then released slowly and the pump valved off.

Before inserting the probe and beginning to sample, the Particulate Data Sheet (Figure IV-29) was checked to see that all the information required was filled out. After a good preliminary leak check was obtained, the initial dry gas meter reading was logged, along with all of

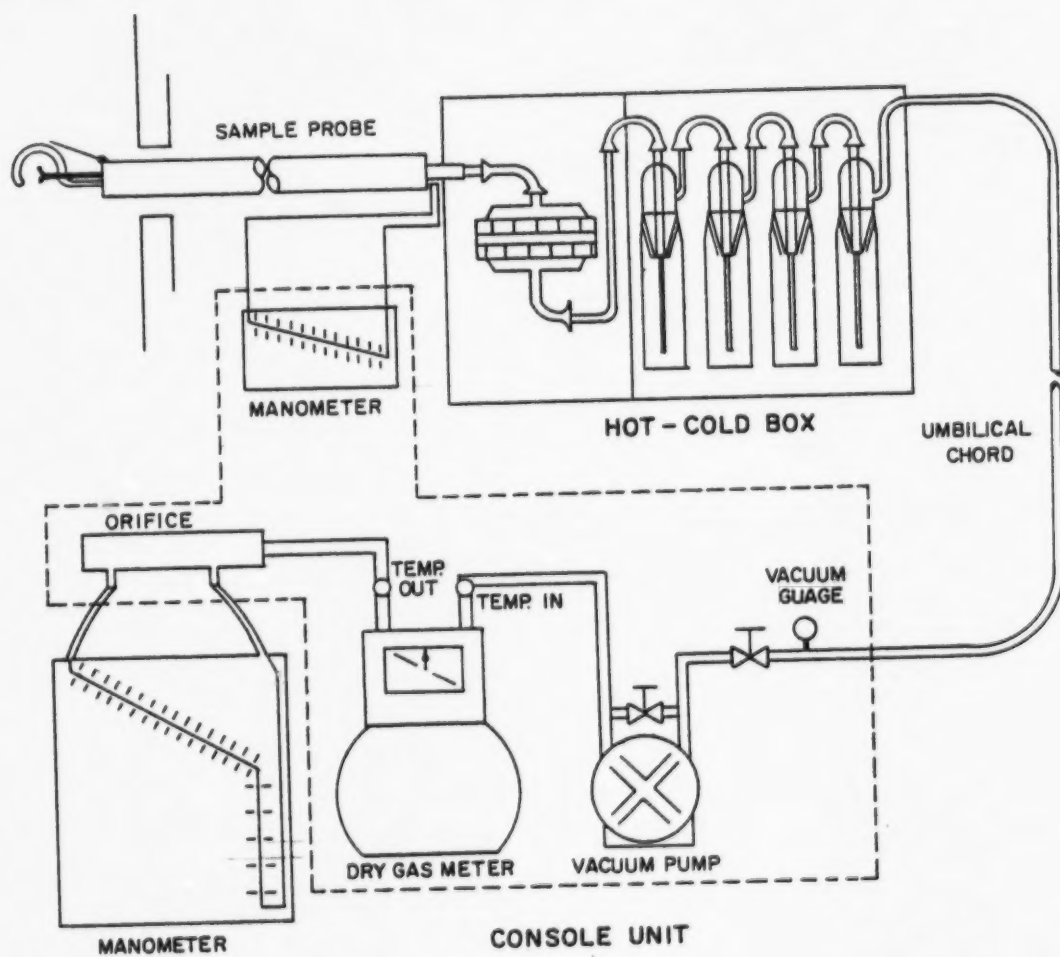


FIGURE IV-28. PARTICULATE SAMPLING TRAIN

SOUTHWESTERN PUBLIC SERVICE COMPANY
PARTICULATE DATA

Plant: _____ Date: _____
Sample Location: _____
Run Number: _____

COMPLETE THIS PART BEFORE SAMPLING

Meter Number: _____ Dry Gas Meter Cal. Factor: _____
Pitot Tube No.: _____ Pitot Tube Cal. Factor: _____
Probe Number: _____ Filter Number: _____
Nozzle Number: _____ Ambient Temp.: _____ °F
Nozzle Diameter: _____ in., cm., mm. (circle one)
Uncorr. Bar. Pressure: _____ in. Hg.
Prelim. Pitot Line Leak Check: o.k., not o.k., (circle one)
Prelim. Probe Leak Check: o.k., not o.k., (circle one)
(less than .02 cfm)
Leak Rate: _____ cf/_____ min.
Static Pressure in Duct: _____ in. H₂O
Operator: Initial when all prelim. checks finished _____

COMPLETE THIS PART AFTER SAMPLING AND BEFORE DISCONNECTING
ANY LINES

Check Cold Box Temp. _____
Check Hot Box Temp. _____
Final Probe Leak Check: o.k., not o.k. (circle one)(less than .02 cfm)
Leak Rate: _____ cf/_____ min.
Final Pitot Line Leak Check: o.k., not o.k. (circle one)

Operator Signature: _____

Time: _____

Figure IV-29. Particulate Data Sheet

the other information required in the Particulate Field Data Form (Figure IV-30). The probe was then inserted into the stack to the inner most point and the pump valve opened. The delta H was immediately adjusted to the isokinetic value indicated by the nomograph. Adjustments to delta H in relation to the delta P were made immediately after moving to a new point. All data was recorded at each point. Close attention was paid to the amount of sampling time at each point. After completing all the points in one port, the pump was valved off and the sampling train moved to the next port. After the last port had been sampled, a post leak check was performed. A leak check was performed before and after disconnecting any portion of the sampling train during a run.

2. Sample Recovery: After completing the post leak check, the impingers were disconnected from the sampling train and capped. The hot box was removed and the glassware capped; the nozzle was removed and capped. The impingers, hot box, and nozzle were then taken to a relatively dust free lab environment for recovery.

The probe was usually washed on the sampling site due to the difficulty in removing it from the site. Any exterior surfaces near the collection end of the probe were wiped off. Any grease on the end that was connected to the glassware was wiped off. The probe liner was then rinsed with analytical reagent grade acetone, catching the washings in the collection bottle. After rinsing, the probe liner was washed using a probe brush and acetone. These washings were added to the collection bottle. The washing procedure was continued until the washings were free of particulate. If there was no visible particulate in the washings, the probe liner was washed at least three times. A separate collection bottle was filled with acetone at each sampling location and labeled "acetone blank". The labels on the collection bottles were filled out completely.

If a probe extension was used between the hot box and cold box and if it was determined that the impinger catch (back half of sampling train) was to be analyzed, the extension was rinsed with water several times. The nozzle was wiped free of exterior particulate and then rinsed with acetone. Next, a nozzle brush and acetone were used to scrub the nozzle clean. All washings were added to the collection bottle.

The glassware in the hot box was disassembled and the particulate caught in the cyclone flask was washed into the collection bottle. The filter holder was taken apart very carefully and the filter, along with the gasket, removed with tweezers. The filter and gasket were placed in the petri dish that the filter was stored in. The petri dish was then sealed with tape. The cyclone, filter holder glassware, fritted disc, and connecting hot box glassware were rinsed with acetone and the washings added to the collection bottle.

The impingers were disconnected and any water or grease on the exterior wiped off. Each impinger was then weighed and this information recorded on the Moisture Fraction Form (Figure IV-31). If it was determined that the impinger catches were analyzed for particulate, they were poured in collection bottles and rinsed with D.I. water. While going through the recovery procedures, all blanks on the recovery form (Figure IV-32) were filled out and the form was signed by all of those involved or in charge of the recovery.

PARTICULATE FIELD DATA

Plant: _____ Sample Location: _____ Date: _____
 Port No.: _____ Start Time: _____ Operator: _____
 Run

Point No.	Time h m s	Dry Gas Meter Reading, cu. ft.	Δ P		Δ H ₂ O		Dry Gas Meter Temp., °F		Vacuum in. Hg	Hot Box Temp., °F	Stack Temp., °F	Cold Box Temp., °F	ISTOPT
			in. H ₂ O	in. H ₂ O	in.	out							
1	8	3	4	5	6	7	8	9	10	11	12	13	14
1	15	16	17	18	19	20	21	22	23	24	25	26	27
1	28	29	30	31	32	33	34	35	36	37	38	39	40
1	41	42	43	44	45	46	47	48	49	50	51	52	53
1	54	55	56	57	58	59	60	61	62	63	64	65	66
1	67	68	69	70	71	72	73	74	75	76	77	78	79
1	80	81	82	83	84	85	86	87	88	89	90	91	92
1	93	94	95	96	97	98	99	100	101	102	103	104	105
1	106	107	108	109	110	111	112	113	114	115	116	117	118
1	119	120	121	122	123	124	125	126	127	128	129	130	131
1	132	133	134	135	136	137	138	139	140	141	142	143	144
1	145	146	147	148	149	150	151	152	153	154	155	156	157
1	158	159	160	161	162	163	164	165	166	167	168	169	170
1	171	172	173	174	175	176	177	178	179	180	181	182	183
1	184	185	186	187	188	189	190	191	192	193	194	195	196
1	197	198	199	200	201	202	203	204	205	206	207	208	209
1	210	211	212	213	214	215	216	217	218	219	220	221	222
1	223	224	225	226	227	228	229	230	231	232	233	234	235
1	236	237	238	239	240	241	242	243	244	245	246	247	248
1	249	250	251	252	253	254	255	256	257	258	259	260	261
1	262	263	264	265	266	267	268	269	270	271	272	273	274
1	275	276	277	278	279	280	281	282	283	284	285	286	287
1	288	289	290	291	292	293	294	295	296	297	298	299	300
1	301	302	303	304	305	306	307	308	309	310	311	312	313
1	314	315	316	317	318	319	320	321	322	323	324	325	326
1	327	328	329	330	331	332	333	334	335	336	337	338	339
1	340	341	342	343	344	345	346	347	348	349	350	351	352
1	353	354	355	356	357	358	359	360	361	362	363	364	365
1	366	367	368	369	370	371	372	373	374	375	376	377	378
1	379	380	381	382	383	384	385	386	387	388	389	390	391
1	392	393	394	395	396	397	398	399	400	401	402	403	404
1	405	406	407	408	409	410	411	412	413	414	415	416	417
1	418	419	420	421	422	423	424	425	426	427	428	429	430
1	431	432	433	434	435	436	437	438	439	440	441	442	443
1	444	445	446	447	448	449	450	451	452	453	454	455	456
1	457	458	459	460	461	462	463	464	465	466	467	468	469
1	470	471	472	473	474	475	476	477	478	479	480	481	482
1	483	484	485	486	487	488	489	490	491	492	493	494	495
1	496	497	498	499	500	501	502	503	504	505	506	507	508
1	509	510	511	512	513	514	515	516	517	518	519	520	521
1	522	523	524	525	526	527	528	529	530	531	532	533	534
1	535	536	537	538	539	540	541	542	543	544	545	546	547
1	548	549	550	551	552	553	554	555	556	557	558	559	560
1	561	562	563	564	565	566	567	568	569	570	571	572	573
1	574	575	576	577	578	579	580	581	582	583	584	585	586
1	587	588	589	590	591	592	593	594	595	596	597	598	599
1	600	601	602	603	604	605	606	607	608	609	610	611	612
1	613	614	615	616	617	618	619	620	621	622	623	624	625
1	626	627	628	629	630	631	632	633	634	635	636	637	638
1	639	640	641	642	643	644	645	646	647	648	649	650	651
1	652	653	654	655	656	657	658	659	660	661	662	663	664
1	665	666	667	668	669	670	671	672	673	674	675	676	677
1	678	679	680	681	682	683	684	685	686	687	688	689	690
1	691	692	693	694	695	696	697	698	699	700	701	702	703
1	704	705	706	707	708	709	710	711	712	713	714	715	716
1	717	718	719	720	721	722	723	724	725	726	727	728	729
1	730	731	732	733	734	735	736	737	738	739	740	741	742
1	743	744	745	746	747	748	749	750	751	752	753	754	755
1	756	757	758	759	760	761	762	763	764	765	766	767	768
1	769	770	771	772	773	774	775	776	777	778	779	780	781
1	782	783	784	785	786	787	788	789	790	791	792	793	794
1	795	796	797	798	799	800	801	802	803	804	805	806	807
1	808	809	810	811	812	813	814	815	816	817	818	819	820
1	821	822	823	824	825	826	827	828	829	830	831	832	833
1	834	835	836	837	838	839	840	841	842	843	844	845	846
1	847	848	849	850	851	852	853	854	855	856	857	858	859
1	860	861	862	863	864	865	866	867	868	869	870	871	872
1	873	874	875	876	877	878	879	880	881	882	883	884	885
1	886	887	888	889	890	891	892	893	894	895	896	897	898
1	899	900	901	902	903	904	905	906	907	908	909	910	911
1	912	913	914	915	916	917	918	919	920	921	922	923	924
1	925	926	927	928	929	930	931	932	933	934	935	936	937
1	938	939	940	941	942	943	944	945	946	947	948	949	950
1	951	952	953	954	955	956	957	958	959	960	961	962	963
1	964	965	966	967	968	969	970	971	972	973	974	975	976
1	977	978	979	980	981	982	983	984	985	986	987	988	989
1	990	991	992	993	994	995	996	997	998	999	1000	1001	1002
1	1003	1004	1005	1006	1007	1008	1009	1010	1011	1012	1013	1014	1015
1	1016	1017	1018	1019	1020	1021	1022	1023	1024	1025	1026	1027	1028
1	1029	1030	1031	1032	1033	1034	1035	1036	1037	1038	1039	1040	1041
1	1042	1043	1044	1045	1046	1047	1048	1049	1050	1051	1052	1053	1054
1	1055	1056	1057	1058	1059	1060	1061	1062	1063	1064	1065	1066	1067
1	1068	1069	1070	1071	1072	1073	1074	1075	1076	1077	1078	1079	1080
1	1081	1082	1083	1084	1085	1086	1087	1088	1089	1090	1091	1092	1093
1	1094	1095	1096	1097	1098	1099	1100	1101	1102	1103	1104	1105	1106
1	1107	1108	1109	1110	1111	1112	1113	1114	1115	1116	1117	1118	1119
1	1120	1121	1122	1123	1124	1125	1126	1127	1128	1129	1130	1131	1132
1	1133	1134	1135	1136	1137	1138	1139	1140	1141	1142	1143	1144	1145
1	1146	1147	1148	1149	1150	1151	1152	1153	1154	1155	1156	1157	1158
1	1159	1160	1161	1162	1163	1164	1165	1166	1167	1168	1169	1170	1171
1	1172	1173	1174	1175	1176	1177	1178	1179	1180	1181	1182	1183	1184
1	1185	1186	1187	1188	1189	1190	1191	1192	1193	1194	1195	1196	1197
1	1198	1199	1200	1201	1202	1203	1204	1205	1206	1207	1208	1209	1210
1	1211	1212	1213	1214	1215	1216	1217	1218	1219	1220	1221	1222	1223
1	1224	1225	1226	1227	1228	1229	1230	1231	1232	1233	1234	1235	1236
1	1237	1238	1239	1240	1241	1242	1243	1244	1245	1246	1247	1248	1249
1	1250	1251	1252	1253	1254	1255	1256	1257	1258	1259	1260	1261	1262
1	1263	1264	1265	1266	1267	1268	1269	1270	1271	1272	1273	1274	1275
1	1276	1277	1278	1279	1280	1281	1282	1283	1284	1285	1286	1287	1288
1	1289	1290	1291	1292	1293	1294	1295	1296	1297	1298	1299	1300	1301
1	1302	1303	1304	1305	1306	1307	1308	1309	1310	1311	1312	1313	1314
1	1315	1316	1317	1318	1319	1320	1321	1322	1323	1324	1325	1326	1327
1	1328	1329	1330	13									

PRELIMINARY MOISTURE DETERMINATION

If preliminary moisture is not determined by the following procedure please state how preliminary moisture was determined:

CONDENSER METHOD

Meter Temp Inlet _____

Meter Temp Outlet _____

Average Temp. °F °R TM = Average temp. + 460 = _____ °R

First Impinger Second Impinger Third Impinger Silica Gel

Final Wt. _____

Initial Wt. _____

Difference _____

Total _____
 Pressure at Meter, PM, _____ inches Hg. (For purposes of this test, PM is equal to atmospheric pressure since ΔH is very small.)

Dry Gas Meter Reading, Final _____

Initial _____

Net Volume _____ ft³

Corrected Volume (V) _____

Dry Gas Meter Calibration Factor, DGMCF _____

$V = (\text{Net Vol}) \times (\text{DGMCF}) = () \times () = \text{_____ ft}^3$

Moisture Fraction =
$$\frac{(\text{MWC}) \times 1.339 \frac{\text{liters}}{\text{gm H}_2\text{O}}}{\left[(\text{MWC}) \times 1.339 \frac{\text{liters}}{\text{gm H}_2\text{O}} \right] + \left[\frac{(V) \times (PM)}{(TM)} \times 501.7 \frac{\text{liters} \cdot \text{°R}}{\text{ft}^3 \text{ in. Hg}} \right]}$$

Moisture Fraction =
$$\frac{() \times 1.339}{\left[() \times 1.339 \right] + \left[() \times () \times 501.7 \right]}$$

Dry Gas Fraction = 1 - (moisture fraction) = 1 - () = _____

When the moisture determination experiment is repeated, use copies of this page for collecting data and recording results. If this is done, calculate the averages:

Average Moisture Fraction _____

Average Dry Gas Fraction _____

FIGURE IV-31. MOISTURE FRACTION FORM

Reproduced from best available copy.



Check List For Recovery of Particulate Samples

Run No. _____

Circle yes or no for each item.

- | | | |
|--|------------|-----------|
| 1. Was the fritted disc broken or chipped? | <u>YES</u> | <u>NO</u> |
| 2. Did the fritted disc contain visible particulate? | <u>YES</u> | <u>NO</u> |
| 3. Was the filter torn or damaged in any way? | <u>YES</u> | <u>NO</u> |
| 4. Was the impinger catch discolored by particulate? | <u>YES</u> | <u>NO</u> |
| 5. Was there condensation or other unusual material
in the cyclone flask? | <u>YES</u> | <u>NO</u> |

Other comments:

Check to see that the following have been done:

1. Levels of containers marked.
2. Labels on containers filled out completely.

Recovered by _____

Figure IV-32. Check List for Recovery of Particulate Samples

3. Sample Analysis: Before the analysis of the particulate catches were done, there had to be a sufficient number of evaporating dishes brought to constant weight. To do this, the dishes were cleaned and then placed in an oven at 105°C and left for 24 hours, then placed in a desiccator and cooled. Each dish was then weighed and placed back in the desiccator. This procedure continued until two consecutive weighings that agreed within 0.0005 grams were obtained. The average of those weighings was taken as the initial weight or tare weight.

The filters and gaskets were treated in the following manner. Using tweezers, the filter was separated from the gasket and folded with the particulate cake inside. It was then placed in an evaporating dish. The gasket was carefully scraped with a scalpel to remove any filter material stuck to it. This material was placed in the evaporating dish with the filter. The gasket was washed off in the petri dish with isopropanol (acetone will dissolve the petri dish). Then the petri dish was washed with isopropanol into an evaporating dish. An equal amount of isopropanol that was used for washing was collected in a dish and used for a blank.

The acetone washings were poured into evaporating dishes. The collection bottles were rinsed out until no more particulate was visible. These evaporating dishes, along with the isopropanol washings, were placed in the evaporation chamber and the fan turned on. After evaporation, the dishes were placed in a desiccator for at least 24 hours.

After desiccation, the dishes were weighed and the procedure for desiccation and weighing was repeated until a constant weight was obtained. The blanks were treated in the same manner as the washings. Blank volumes should correspond to the volumes used for washing; if not, the exact volumes used were recorded. Results of the weighings were recorded on the appropriate forms for submission. (See Figures IV-33 and IV-34.)

4. Reagents for Particulate:

D.I. Water (as furnished by System Lab).

Acetone (reagent grade, $\geq$ 0.001 percent residue, in glass bottles).

Isopropanol (reagent grade).

Filters (Reeve Angel 900 AF glass fiber).

F. Calibration and Standardization For Particulate

1. Acetone Residual Purity Check: Each lot of acetone used in recovery was checked for purity before use. Results were recorded on the Acetone Purity Check Form, Figure IV-35. To do this, 500 ml of acetone was poured into a tared evaporating dish and evaporated. The weight gain was obtained after desiccation. The weight gain should not be greater than 0.001 percent of the weight of acetone used. For a 500 ml sample, this would be no more than about 3.93 mg weight gain. The exact amount would depend on the density of the acetone in the lot.

2. Dry Gas Meter Calibration: An initial and final calibration was done on the dry gas meter. The procedure used for the calibration of the dry gas meters can be found in the Calibrations Section.

Particulate Results

Plant _____ Particulate Run # _____
 Location _____ Date Sampled _____
 Purpose _____ Date Recovered _____
 Filter # _____

	Evap. Dish #	Final Weight	Tare Weight	Weight Gain
Acetone Blank				
Isopropanol Blank				
Total Blank Weight				

	Evap. Dish #	Final Weight	Tare Weight	Weight Gain
Filter + Evap. Dish				
Filter Washings				
Probe & Glassware				
Washings				
Total Weight Gain				

(Total Weight Gain)-(Total Blank Weight) = (Net Weight Gain)
 _____ - _____ = _____

FIGURE
IV-33
PARTICU-
LATE
BLANK
RESULTS

PARTICULATE RESULTS

Plant _____

Particulate Run # _____

Location _____

Date Sample _____

Purpose _____

Date Recovered _____

Filter # _____

	Evap. Dish #	Final Weight	Tare Weight	Weight Gain
Filter + Evap. Dish				
Filter Washings				
Probe & Before Filter Glassware Washings				
Impinger & After Filter Glassware Washings				
Total Weight Gain				

(Total Weight Gain) - (Total Blank Weight) = (Net Weight Gain)

_____ - _____ = _____

FIGURE IV-34. PARTICULATE RESULTS

Date Acetone Received _____
 Lot # _____
 Checked by _____
 Date Checked _____

Date Checked _____

[illegible]

3. Orifice Calibration: The calibration of the orifice was done before each test. The procedure to be used for the calibration of the orifice can be found in the Calibrations Section.

4. Magnehelic Calibrations: Any magnehelic used for the measurement of pressure differentials was calibrated against incline manometers. The procedure used for these calibrations can be found in the Calibrations Section.

5. Reporting of Results: The results of the analysis were reported to Engineering Services as net weight gain in grams. This value was calculated according to the following formula:

$$\text{Total Weight Gain} - \text{Total Blank Weight} = \text{Net Weight Gain} \quad (10)$$

The total blank weight must be adjusted by using a multiplication factor if the volume of the blank was not the same as the sample volume. All other calculations were done by Engineering Services.

6. Quality Assurance: As a check on the quality of the data collected, the standard deviation of the PMR (Particulate Mass Emission) for the three runs was calculated along with the 90 percent confidence limits. The formulas for these calculations are as follows:

$$S \text{ PMR} = \frac{(\text{PMR}_1 - \overline{\text{PMR}})^2 + (\text{PMR}_2 - \overline{\text{PMR}})^2 + (\text{PMR}_3 - \overline{\text{PMR}})^2}{2}^{1/2} \quad (11)$$

where

$S \text{ PMR}$ = The calculated standard deviation for the three sample runs

PMR_1 ,

PMR_2 ,

PMR_3 = Particulate Mass Emission rate for sample run 1, 2, and 3

$\overline{\text{PMR}}$ = Average particulate mass emission rate of the three sample runs.

2 = Number of replicates minus one.

$$90\% \text{ confidence limit} = \overline{\text{PMR}} \pm (2.92) (S \text{ PMR}) / \sqrt{n} \quad (12)$$

where

$\overline{\text{PMR}}$ = The average of the three replicates

S PMR = Estimated standard deviation of PMR based on three replicates

2.92 = 95th percentile of the student t - distribution with 2 degrees of freedom

n = The number of replicates, i.e., n = 3 for this case.

Example:

For a given field test, $\overline{\text{PMR}} = 22.86 \text{ g/hr.}$ and $S \left\{ \text{PMR} \right\} 1.32 \text{ g/hr.}$

The reported value with 90% confidence limits would be

$$22.86 \text{ g/hr.} \pm (2.92) (1.32 \text{ g/hr.}) / 3 \quad (13)$$

or

$\overline{\overline{\text{PMR}}}$ (grand mean particulate mass emission) is

$$20.6 \text{ g/hr.} \leq \overline{\overline{\text{PMR}}} \leq 25.1 \text{ g/hr.}$$

G. Collection and Analysis For SO₂

1. Sample Collection: The sampling train was set up with five midjet impingers. The first impinger contained 15 ml of 80 percent isopropanol. The second and third impingers contained 15 ml of 3 percent hydrogen peroxide. The fourth impinger was empty and the fifth contained silica gel. The train was assembled as shown in Figure IV-36, making sure that the connecting glassware was on the impingers in the correct sequence. Crushed ice was placed around the impingers. A collection bottle was filled with some of the H₂O₂ and labeled "H₂O₂ Blank."

A preliminary and a post leak check performed in the following manner. After the sampling train was completely assembled, the probe tip was plugged with a rubber stopper and the control valve to the pump opened. A vacuum of at least 5 inches of Hg was pulled. The needle on the dry gas meter was observed for any movement after the vacuum was reached. A leak rate not in excess of two percent of the average sampling rate was acceptable; however, if the rate was not zero, all connections were checked. The plug was then released slowly and the pump valved off.

After a good preliminary leak check was obtained, the initial dry gas meter reading was logged, along with all of the other information

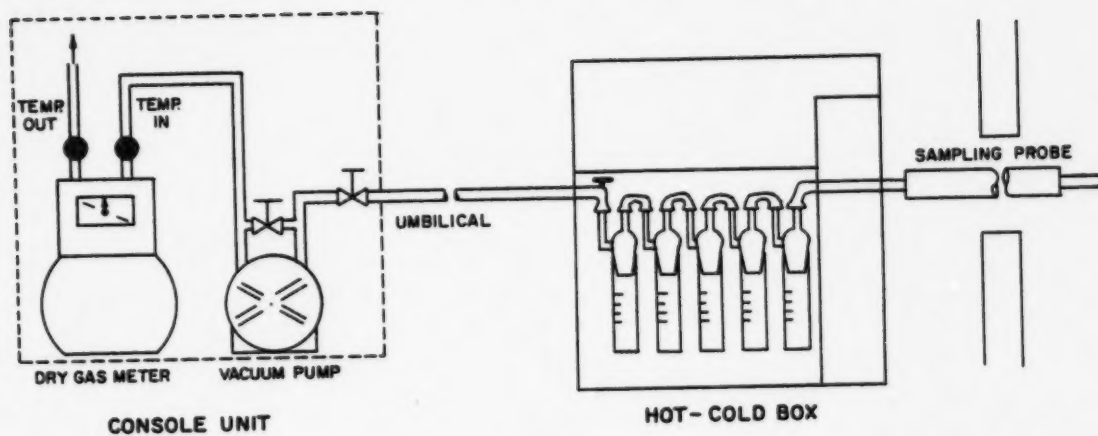


FIGURE IV-36. SO_2 SAMPLING TRAIN

requested on the SO₂ Sample Field Data Form (Figure IV-37). The probe was then inserted into the stack and the pump control valve opened and adjusted to the pre-determined sampling rate. Readings were logged every five minutes during sampling. At the conclusion of the run, the pump control valve was closed, the readings recorded, and a post leak check conducted in the same manner as described on the preceding page. After the leak check had been completed, the probe from the impinger train was disconnected and the train was purged with clean ambient air for 15 minutes.

2. Sample Recovery: After purging for 15 minutes, the connecting glassware was disconnected and laid aside. The number of ml in each impinger was noted and recorded. The contents of impinger #1 were poured into a collection bottle and labeled "isopropanol." Contents of the #2, #3, and #4 impingers were poured into a separate collection bottle. These impingers and their connecting glassware were washed with D.I. water and these washings added to the collection bottle. The container level was marked and the label (Figure V-38) was completely filled out after completing the recovery procedure. A collection bottle was filled with D.I. water and labeled "D.I. Water Blank." The containers were then sealed with tape if there was going to be an extended period of time between collection and analysis. A recovery check list was provided for each run and was filled out completely and signed at the conclusion of the recovery procedures. (See Figure IV-39.)

3. Sample Analysis: The level of the liquid in the collection bottle was noted and if there was any loss, it was indicated on the analysis form (Figure IV-40). The contents of the bottle were transferred to a volumetric flask big enough to allow for the dilution of each sample to a constant volume. A 20-ml aliquot or an aliquot of sufficient volume to require at least 5 ml of 0.01N barium perchlorate solution for titration was pipetted into an Erlenmeyer flask. Eighty ml of 100 percent isopropanol and four drops of Thorin indicator were added. The sample was titrated with 0.0100N barium perchlorate to the salmon pink endpoint. The titration was repeated on a second aliquot. The replicate titrations must agree within 1 percent or 0.2 ml, whichever is larger. If the replicate titrations agreed according to the limits, the two numbers were averaged and used for V_t in the formula. V_a was the volume of the aliquot. V_{soln} was the constant volume that each sample was diluted to. Results were recorded on the SO₂ Data Summary Form (Figure IV-41).

A blank was run with each set of analysis. The blank consisted of a volume of deionized water and H₂O₂, the same as the aliquot used and 80 ml of 100 percent isopropanol.² A set of standards was also run with each set of analysis for quality assurance. The preparation of these standards is described under Reagents. The treatment of the quality assurance data is covered under Quality Assurance.

4. Reagents for SO₂:

80 percent AR Grade Isopropanol prepared from 100 percent isopropanol.

3 percent H₂O₂ (prepared daily, diluted 10 ml of the 30 percent solution to 100 ml with D.I. water).

SO₂ SAMPLE FIELD DATA

EPA METHOD 6

PLANT/LOCATION _____ DATE _____

PURPOSE _____ OPERATOR _____

Pbar _____ in. Hg. METER NO. _____ D.G.M.C.F. _____

SAMPLE NO. _____ PRE LEAK CK. _____ POST LEAK CK. _____

PORT	TIME	DRY GAS MTR. READING	ΔP in. Hg	ΔH in. Hg	DRY GAS MTR. TEMPERATURE		VAC in. Hg	HOT BOX TEMP. °F	COLD BOX TEMP. °F	STACK TEMP. °F
					IN	OUT				

TOTAL SAMPLE TIME _____ MIN. _____ X _____ = _____ ft.³
 Net Vol. D.G.M.C.F. Corrected Vol. at
 Meter Cond.

SAMPLE NO. _____ PRE LEAK CK. _____ POST LEAK CK. _____

PORT	TIME	DRY GAS MTR. READING	ΔP in. Hg	ΔH in. Hg	DRY GAS MTR. TEMPERATURE		VAC in. Hg	HOT BOX TEMP. °F	COLD BOX TEMP. °F	STACK TEMP. °F
					IN	OUT				

TOTAL SAMPLE TIME _____ MIN. _____ X _____ = _____ ft.³
 Net Vol. D.G.M.C.F. Corrected Vol. at
 Meter Cond.

FIGURE IV-37. SO₂ SAMPLE FIELD DATA FORM

Plant _____
Location _____
Sample # _____
Date _____
Recovered by _____

Figure IV-38. Bottle Label

Check List For Recovery of SO₂ Samples

Run No. _____

1. Indicate the approximate volume of liquid in each impinger to the nearest ml.

				contains silica gel
_____	_____	_____	_____	_____
#1	#2	#3	#4	#5

2. Indicate if there was any discoloration in the impinger liquid by circling either yes or none.

<u>yes/none</u>	<u>yes/none</u>	<u>yes/none</u>	<u>yes/none</u>	<u>yes/none</u>
_____	_____	_____	_____	_____
#1	#2	#3	#4	#5

3. indicate the extent that the silica gel was spent by circling one of the following:

slightly pink localized pink almost entirely pink

Other comments:

Recovered by _____

Check to see that the following have been done:

1. Levels of containers marked.
2. Lables on containers filled out completely.
3. Container sealed for shipment.
4. H₂O₂ and D.I. water blanks collected.

Figure IV-39. Check List for Recovery of SO₂ Samples

SO₂ ANALYSIS
EPA METHOD 6

Plant Name: _____

Titrant: _____

Purpose: _____

Normality of Titrant: _____

Date Sampled: _____

Was there any loss of sample during

Date Analyzed: _____

transport? Yes _____ No _____

Analyst: _____

If yes, which containers: _____

Sample No.	Ml of Titrant Used			Total Solution Volume	Volume of Aliquot
	1	2	Avg.		

FIGURE IV-40. SO₂ ANALYSIS FORM

FIGURE IV-41. SO₂ DATA SUMMARY FORM

Plant Name: _____
Location: _____
Purpose: _____
Date Sampled: _____
Date Analyzed: _____

[illegible]

Primary standard KHP (for standardization of NaOH): dried about 3 grams of KHP ($\text{KHC}_8\text{H}_4\text{O}_4$) at 110°C for one hour and then cooled for at least 30 minutes in a desiccator. Weighed out and dissolved 0.025 grams, 0.050 grams, and 0.100 grams of the standard KHP in 50 ml of D.I. water.

0.01 N NaOH solution (for standardization of H_2SO_4): dissolved 0.40 grams of NaOH in 300 ml of D.I. water and diluted to one liter.

0.01N H_2SO_4 (for standardization of $\text{Ba}(\text{ClO}_4)_2$): added 3 ml of concentrated H_2SO_4 to 500 ml of D.I. water and diluted to one liter. Took 100 ml of this solution and diluted to one liter.

0.01N barium perchlorate solution: dissolved 1.95 grams of barium perchlorate [$\text{Ba}(\text{ClO}_4)_2 \cdot 3\text{H}_2\text{O}$] in 200 ml of D.I. water and diluted to one liter with A.R. grade isopropanol.

Phenolphthalein Indicator: used as prepared by the System Lab.

Thorin Indicator: dissolved 0.20 grams of Thorin Indicator in 100 ml of D.I. water.

10 Percent Potassium Iodide Solution: dissolved 10 grams of KI in D.I. water and diluted to 100 ml.

5. Standard Quality Assurance Samples: All standards were made from dilutions of a 1N solution of H_2SO_4 . For the 1N H_2SO_4 solution, 26.7 ml of concentrated H_2SO_4 was diluted (specific gravity of 1.84) to one liter. The 1N solution was standardized in the same manner that the H_2SO_4 solution used for barium perchlorate standardization was done. The weight of KHP was 2.0000 g, 1.0000 g, and 0.4800 g. After the 1N solution of H_2SO_4 had been standardized, it was used in the following manner for each standard:

80 Percent Range Standard (0.0048N H_2SO_4): added 4.8 ml of 1N H_2SO_4 solution to a 1-liter volumetric flask and diluted to the line.

Median Range Standard (0.0028N H_2SO_4): added 2-8 ml of 1N H_2SO_4 solution to a 1-liter volumetric flask and diluted to the line.

20% Range Standard (0.0012N H_2SO_4): added 1.2 ml of 1N H_2SO_4 solution to a 1-liter volumetric flask and diluted to the line.

These standards were approximately equivalent to the following amounts of SO_4 :

80 Percent Range Standard	-	0.235 mg/ml
Median Range Standard	-	0.137 mg/ml
20 Percent Range Standard	-	0.059 mg/ml

H. Calibrations and Standardization for SO_2

1. Isopropanol Purity Checks: Each lot of isopropanol was checked for peroxide impurities. To do this, 10 ml of isopropanol was shaken with 10 ml of freshly prepared 10 percent KI. A blank was prepared

using 10 ml of D.I. water. After one minute, the absorbance was read at 352 nanometers on a spectrophotometer. The absorbance must not exceed 0.1. A form for recording this procedure was provided (see Figure IV-42).

2. Dry Gas Meter Calibration: An initial and final calibration was done on the dry gas meter. The procedure used for the calibration of the dry gas meters can be found under Calibrations in this report. The initial and final calibration factors must not deviate by more than 5 percent. If they differ by more than 5 percent, the calibration factor that would yield the lower gas volume for each test run was used.

3. Standardization of Barium Perchlorate: the standardization for the barium perchlorate titrant was performed using a standard solution of H_2SO_4 which is standardized using a standard solution of NaOH which is standardized using primary standard KHP. Results were recorded on Figure IV-43.

4. Standardization of NaOH (used form shown in Figure IV-44):

Prepared the three KHP solutions and the NaOH solution as described in the Reagent Section of these procedures.

Added two or three drops of phenolphthalein to the first KHP solution and titrated with the NaOH until the pink endpoint was reached.

Recorded the number of ml required for each sample and calculated the normality of the NaOH using the following formula:

$$N = \frac{M}{(V_t) (204.22)} \quad (14)$$

where

N = Normality of the NaOH

M = mg of KHP used

V_t = Volume of NaOH solution used (ml)

204.22 = Equivalent weight of KHP.

Averaged the normalities calculated for all three samples and assigned this value to the solution. Put the normality and date of standardization on the container.

5. Standardization of H_2SO_4 (used form shown in Figure IV-45):

Prepared the H_2SO_4 solution as described in the Reagent Section of these procedures.

Titrated two or three 20 ml samples of H_2SO_4 with the standard NaOH solution to the faint pink endpoint.

Calculated the normality of the H_2SO_4 solution using the following equation:

$$N_2 = \frac{N_1 V_1}{V_2} \quad (15)$$

Isopropanol Check for Peroxide Impurity

Lot # _____

Absorbance

Blank

Sample #1

Sample #2

Sample #3

Absorbance Must Be 0.1

Figure IV-42. Isopropanol Check for Peroxide Impurity

Standardization of Barium Perchlorate

Date _____

Solution I.D. # _____

Analyst _____

Normality of H_2SO_4 _____

Sample #	ml of Standard H_2SO_4 Used	ml of $Ba(ClO_4)_2$ Titrant Used
----------	----------------------------------	-------------------------------------

Calculations:

$$N_2 = \frac{N_1 V_1}{V_2}$$

Where

N_2 = Normality of $Ba(ClO_4)_2$

N_1 = Normality of H_2SO_4

V_1 = Volume of H_2SO_4 titrated

V_2 = Volume of $Ba(ClO_4)_2$ titrant used

Sample #	Normality of $Ba(ClO_4)_2$
----------	----------------------------

Average

Figure IV-43. Standardization of Barium Perchlorate

Standardization of NaOH

Date _____

Solution I.D. # _____

Analyst _____

Sample #	mg of KHP Used	ml of NaOH Titrant
----------	----------------	--------------------

Calculations:

$$N = \frac{m}{(V_t)(204.22)}$$

Where

N_2 = Normality of NaOH

m = mg of KHP used

V_t = Volume of NaOH titrant used

204.22 = Equivalent weight of KHP

Sample #	Normality of NaOH
----------	-------------------

Average

Figure IV-44. Standardization of NaOH

Standardization of H_2SO_4

Date _____

Solution I.D. # _____

Analyst _____

Normality of NaOH _____

Sample #	ml of H_2SO_4	ml of NaOH
	Solution Titrated	Titrant Used

Calculations:

$$N_2 = \frac{N_1 V_1}{V_2}$$

where

N_2 = Normality of H_2SO_4

N_1 = Normality of the NaOH solution

V_1 = Volume of NaOH titrant used

V_2 = Volume of H_2SO_4 titrated

Sample #	Normality of H_2SO_4
----------	--------------------------------------

Average

Figure IV-45. Standardization of H_2SO_4

where

N_2 = Normality of the H_2SO_4
 N_1 = Normality of the NaOH solution
 V_1 = Volume of NaOH titrant used (ml)
 V_2 = Volume of H_2SO_4 titrated.

Averaged the three normalities and assigned that value to the H_2SO_4 solution. Put the normality and date of standardization on the container.

The standard H_2SO_4 solution was used repeatedly once it had been standardized. However, if the date on the bottle indicated that it was more than six months old, it was discarded and a new solution standardized.

6. Standardization of the Barium Perchlorate Solution: Prepared the barium perchlorate as described in the Reagent Section of these procedures.

Prepared three samples containing 25 ml of standard H_2SO_4 and 100 ml of isopropanol.

Added four drops of Thorin indicator and titrated with $Ba(ClO_4)_2$ to the endpoint.

Calculated the normality of $Ba(ClO_4)_2$ by using the following formula:

$$N_2 = \frac{N_1 V_1}{V_2} \quad (16)$$

where

N_2 = Normality of $Ba(ClO_4)_2$
 N_1 = Normality of H_2SO_4 solution
 V_1 = Volume of H_2SO_4 solution titrated
 V_2 = Volume of $Ba(ClO_4)_2$ titrant used.

Averaged the normalities from all three titrations and assigned that normality to the $Ba(ClO_4)_2$ solution. Labeled the bottle with the normality and date of standardization.

The level left in the H_2SO_4 bottle and the $Ba(ClO_4)_2$ bottle was marked each time an amount was used. All precautions were taken to prevent the evaporation of the solutions after standardization.

I. Calculations of Final Results

The System Lab used a computer program from its file which calculated the dry gas volume at standard conditions (V_{mstd}); concentration of SO_2 in lb/dscf (C_{SO_2}); concentration of SO_2 in ppm; and concentration in $lb/10^6$ Btu. A file copy of the program can be found in Table IV-1.

The program used the following formulas to calculate the results.

```

0001      READ (5,100,END=999) F, NSAMP
0002      100 FORMAT (F6.0,12)
0003      WRITE(6,104)
0004      103 FORMAT(1H //13X, 'VMSTD',8X,'CS02',9X,'PPM',
&6X, 'LB/10**6BTU',4X,'RUN'
&/13X,'(DSCF)',3X,'(LB/DSCF)',32X,'NO.')
```

0005

```

      104 FORMAT(1H1,25X,'SOUTHWESTERN PUBLIC SERVICE CO.',
&/35X,'SYSTEM LAB',
&/33X'SULFUR DIOXIDE')
      WRITE (6,103)
      2 CONTINUE
      TCS02 = 0.
      TPPM = 0.
      E = 0.
      DO 3 I=1,NSAMP
      READ (5,101) J,VM,PBAR,TM,VT,VTB,AN,VSOLN,VA,02
      101 FORMAT(A5,F6.3,F6.2,F5.0,F5.2,F5.2,F5.2,F5.0,F4.0,F4.1)
      VMSTD = 17.65 * VM * PBAR /TM
      CS02=((7.061E-5)*(VT-VTB)*AN*(VSOLN/VA))/VMSTD
      PPM = CS02 /1.659413E-7
      E = CS02 * F * 20.9 /(20.9 - 02)
      WRITE (6,102) VMSTD,CS02,PPM,E,J,
      102 FORMAT(//10X,F8.2,2X,F10.7,4X,F8.2,6X,F8.6,5X,A5)
      TCS02 = TCS02 + CS02
      TPPM = TPPM + PPM
      TE = TE + E
      3 CONTINUE
      ACS02 = TCS02 / NSAMP
      APPM = TPPM / NSAMP
      AE = TE / NSAMP
      WRITE(6,106)
      106 FORMAT(1H ,20X,'-----',4X,'-----',
&5X,'-----')
      WRITE(6,107)
      107 FORMAT(1H ,1X,'AVERAGE')
      WRITE(6,108)ACS02,APPM,AE
      108 FORMAT(1H ,19X,F10.7,4X,F8.2,6X,F8.6)
      READ(5,100,END=999)F,NSAMP
      IF(F.NE.0.) GO TO 2
      WRITE(6,105)
      105 FORMAT(1H-,//40X,'-----',
&/40X,'PREPARED BY')
```

0037

```

      999 STOP
      END
0038
```

TABLE IV-1. DRY GAS VOLUME COMPUTER PROGRAM

SO₂ Calculations

$$V_m(\text{std}) = 17.65 (\text{dgmcf}) \frac{(V_m P_{\text{bar}})}{T_m} \quad (17)$$

where

$V_m(\text{std})$ = dry gas volume measured by dry gas meter (standard conditions) (dscf)

$$17.65 = \frac{\text{std. temp.}}{\text{std. press.}} = \frac{528^\circ \text{ R}}{29.92 \text{ in Hg}}$$

dgmcf = dry gas meter correction factor

V_m = dry gas volume as measured by dry gas meter (dcf) without (dgmcf)

T_m = Avg. dry gas meter absolute temperature ($^\circ\text{R}$)

P_{bar} = barometric pressure at dry gas meter orifice (in. Hg).

$$C_{\text{SO}_2} = \frac{(7.061 * 10^{-5}) (V_t - V_{\text{tb}}) N \left(\frac{V_{\text{soln}}}{V_a} \right)}{V_m(\text{std})} \quad (18)$$

where

C_{SO_2} = concentration of sulfur dioxide, dry basis, standard conditions (lb/dscf)

$7.061 * 10^{-5}$ = conversion factor (lb/meq)

V_t = volume of barium perchlorate titrant used for sample (ml)

V_{tb} = volume of barium perchlorate titrant used for blank (ml)

N = normality of barium perchlorate titrant (meq/ml)

V_{soln} = total solution volume of sulfur dioxide sample (ml)

V_a = volume of sample aliquot titrated (ml).

To convert C_{SO_2} in lb/dscf to ppm:

$$\text{ppm} = \frac{C_{\text{SO}_2}}{(2.59 * 10^{-9}) (64.07)} \quad (19)$$

$$E = (C) (F) \frac{20.9}{20.9 - \text{percent O}_2} \quad (20)$$

where

E = concentration in lb/10⁶ BTU
C = concentration in lb/dscf
F = factor specific for type of fuel being burned
percent O₂ = O₂ as measured by Orsat
20.9 = concentration of O₂ in air.

J. Quality Assurance

Four control charts were maintained during the special testing program to assure the quality of the data reported for SO₂ determinations. The charts for the three standard solutions were run with each batch of samples. The mg of SO₂ present (as determined by the titration procedure) was used to figure the percent difference in the measured value from the true value. This percent difference was plotted versus the date of analysis making up the control chart. (See Figure IV-46.) The formulas for calculation of mg of SO₂ and percent difference are as follows:

$$m = (N) (V_t) (49.04) \quad (21)$$

where

m = mg of H₂SO₄ present
N = normality of barium perchlorate titrant
V_t = ml of barium perchlorate titrant used
49.04 = equivalent weight of H₂SO₄.

$$d = \frac{C_m - C_t}{C_t} \times 100 \quad (22)$$

where

d = % difference
C_m = measured concentration
C_t = true concentration.

Comments were placed on the control chart when a new batch of standards was made up or a new reagent was used. The fourth control chart was a range chart for the Ba(ClO₄)₂ normality. The normality of the Ba(ClO₄)₂ solution being used was checked periodically and the difference from the original value calculated. This percent difference was plotted in the same manner as above. Interpretation of the quality control charts is discussed under Quality Control and Assurance.

K. Collection and Analysis For NO_x

1. Preparation of NO_x Flasks Prior To Sampling: Flasks and stopcocks were rinsed with^x deionized (DI) water. The stopcock was carefully checked to make sure the sleeve was in good shape and intact. Excess water was drained from the flask and stopcock. Stopcock glass joints were slightly greased with a chlorofluorocarbon-based stopcock grease, (e.g., Halocarbon grease).

SO₂ QUALITY ASSURANCE SAMPLES

Date Analyzed _____

Titrant _____

Analyst _____

Titrant Solution I. D. # _____

Normality of Titrant _____

Sample No.	Ml of Titrant Used			Volume of Aliquot	True Mg	Calculated Mg	O/O Difference
	1	2	Avg.				

FIGURE IV-46. SO₂ QUALITY ASSURANCE SAMPLES

2. Sample Collection: Barometric pressure (uncorrected) was recorded. Twenty-five ml of freshly prepared NO_x absorbing solution was pipeted into a sampling flask. A sufficient quantity of absorbing solution was retained for preparing at least three standards and a blank for each twelve samples collected, i.e., at least 125 ml for each twelve samples.

The sampling probe was placed at a point between one-third and two-thirds into the stack. The sample train was assembled as shown in Figure IV-47. It is advisable to make the connections between the probe and flask as short as possible. The procedure outlined below was then followed:

Zeroed the manometer.

With manometer valve shut and pump valve open (Figure IV-48) and stopcock in P position (Figure IV-48), pump started.

Shut pump valve and opened manometer valve, then set stopcock in E position (Figure IV-48).

Opened pump valve and evacuated flask to 3 inches Hg absolute pressure or less (approximately 24 inches on mercury manometer).

Shut pump valve and checked manometer for fluctuation in mercury level (this leak checked the sample train between manometer and flask). If the fluctuation was greater than 0.4 inches Hg per minute, the leak rate was unacceptable and was changed.

If leak rate was zero or acceptable, initial temperature (T_i) and pressure (P_i) of the flask was recorded and stopcock set to P position. If the time between leak check and sampling was greater than one minute, then the flask pressure was re-checked just prior to sampling.

Pump valve opened and sample train purged. If condensation in the sample line was evident during purging, then the line between the stack and flask was heated by using a blow dryer.

Pump valve shut and when ready to sample, stopcock set in S position (Figure IV-48); sample pulled for approximately 20 seconds.

Stopcock set in P position, being extremely careful not to disengage stopcock from flask; flask disconnected from the sampling train and shaken continuously for 5 minutes.

3. Sample Recovery: After allowing the sample flasks to set for a minimum of 16 hours, each flask was shaken for at least 2 minutes. Barometric pressure was recorded (P_{bar}), as well as each flask temperature (T_f) and pressure (P_f), using the NO_x Sampling Data Form in Figure IV-49. Flask contents were transferred to a collection bottle and the flask and stopcock were rinsed with at least two 10-ml portions of DI water. The rinse water was added to the collection bottle. A blank, consisting of 25 ml NO_x absorbing solution and approximately the same amount of DI water used for rinsing each flask and stopcock was prepared. To each collection bottle, 1N sodium hydroxide (30-50 drops) was added until just basic to litmus paper (pH approximately 9-12) and then was checked by dipping a clean stirring rod into the solution and touching it to the litmus or pH paper. The collection bottle level was marked to determine any leakage between time of sample recovery and analysis. At least 100 ml of NO_x absorbing solution was retained for each twelve samples that were recovered. This was used in preparing NO_x standards during analysis. Each container was labeled to clearly identify its contents. Finally, each container was sealed with plastic tape.

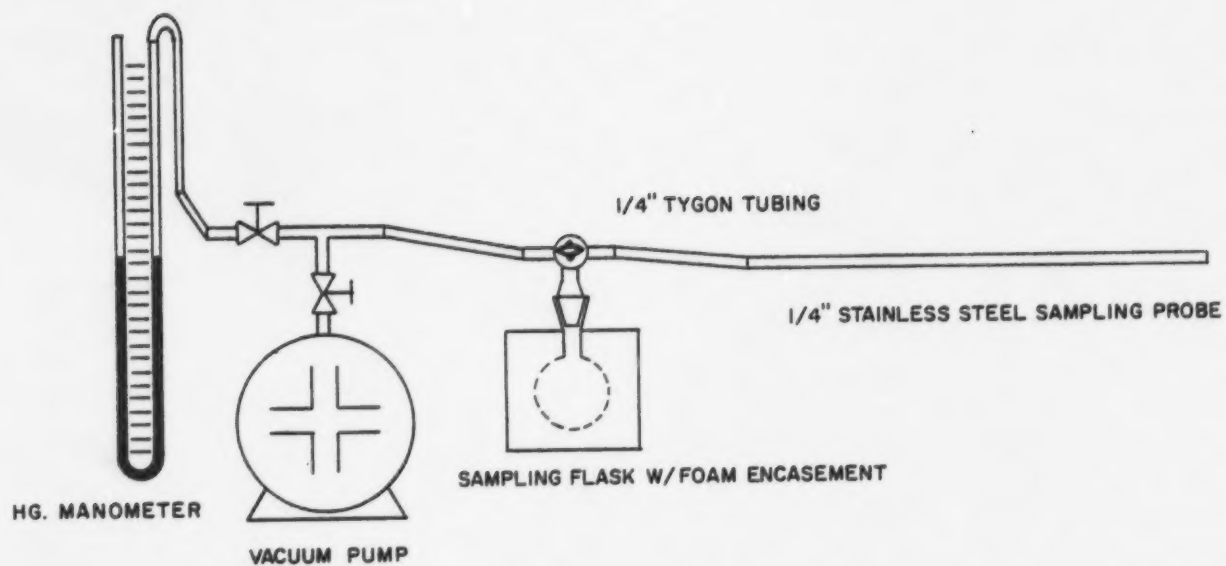


FIGURE IV-47. NO_x SAMPLING TRAIN



FIGURE IV-48. FLASK STOPCOCK POSITIONS

NO_x SAMPLING DATA
EPA METHOD 7

Plant/Location _____ Sample Date _____

Purpose _____

Pbar Uncorrected _____ in. Hg Temp. _____ °F

Sample #	Time	Flask No.	Leak Rate	Pi in. Hg	Ti F	Pf in. Hg	Tf F	Comments	Orast
									Leak Ck
									Time
									CO ₂
									O ₂
									Leak Ck.
									Time
									CO ₂
									O ₂
									Leak Ck.
									Time
									CO ₂
									O ₂

Sample Recovery Date _____

Pbar Uncorrected _____ in. Hg Temp. _____ °F

Sampled by _____

Recovered by _____

Check to see that the following have been done at the time of recovery.

- ☐ 1. Levels marked on containers
- ☐ 2. Labels on containers filled out completely
- ☐ 3. Container sealed for shipment

FIGURE IV-49. NO_x SAMPLING DATA FORM

4. Sample Analysis: The level in each collection bottle was noted and an indication was made on the analysis sheet (Figure IV-50) if any sample was lost during shipment and storage. Three standards were prepared to be run with each set of analyses. A 500 ug (low range), 1000 ug (mid range), and 1750 ug (high range) NO_2 standard was prepared in the following manner:

500 ug NO_2 standard	1000 ug NO_2 standard	1750 ug NO_2 standard
Added 20 ml KNO_3 "working standard solution" to an evaporating dish. . .	Added 40 ml KNO_3 "working standard solution" to an evaporating dish. . .	Added 80 ml KNO_3 "working standard solution" to an evaporating dish. . .

. . . then pipeted 25 ml NO_x absorbing solution into each evaporating dish." The standard was diluted with deionized water to the same volume as each sample collection bottle and then added 1N sodium hydroxide, dropwise, until each standard solution was just basic to litmus (pH approximately 9-12).

The contents of each sample collection bottle was transferred into an evaporating dish and each bottle was rinsed twice with 5 ml portions of DI water. The following steps were then taken.

Evaporated each solution, including freshly prepared standards, to dryness over a steam bath; removed, and allowed to cool.

Added 2 ml phenoldisulfonic acid to each and tritrated thoroughly with a glass rod or polyethylene policeman. (Made sure acid contacted all of residue.)

Added 1 ml DI water and four drops of concentrated sulfuric acid.

Heated each solution over a steam bath for three minutes with occasional stirring; removed, and allowed to cool.

Added 10 ml DI water and mixed well by stirring.

Added 15 ml of fresh, concentrated ammonium hydroxide dropwise to excess (pH approximately 10.0). Checked pH of at least one sample with pHydron paper to assure that a pH of 10.0 is being attained.

Filtered each solution into a 250 ml volumetric flask. Rinsed each evaporating dish thoroughly with DI water, also filtering each rinse. Washed all color completely from each filter into the corresponding 250 ml volumetric flask and diluted to the mark with DI water.

Mixed each flask solution well.

Read and recorded on analysis sheet the percent transmittance of each sample against a blank at 420 nm on a spectrophotometer.

Converted the photometric reading to ug NO_2 by means of the calibration curve and recorded on analysis sheet.

L. Calculation of Final Results

The System Lab used a computer program from its file to calculate the dry gas volume at standard conditions (V_{sc}); concentration of NO_x in lb/dscf (C); concentration of NO_x in ppm_{dry}; and concentration in lb/10⁶ Btu. A file copy of the program can be found in Table IV-II.

The program uses the following formulas to calculate the results.

FIGURE IV-50. NO_x ANALYSIS FORM

Date _____

Plant _____ Sample Recovery Date _____

Purpose _____

Pbar Uncorrected _____ in. Hg Temp. _____ °F

Sample Date _____

Bar Uncorrected _____ in. Hg Temp. _____ °F

Phar Corrected _____ in. Hg

[illegible]

*Pressure corrected for Temp. and Elevation


```

0001      DIMENSION PLANT(4),SAMDAT(4),ITEMP(2),SAMLO(3),
          &TITLE(4,4)
0002      DATA BLANK/'  '/
0003      30 CONTINUE
0004      READ (5,100,END=999) F,NSAMP
0005      100 FORMAT (8X,F5.0,7X,I2)
0006      WRITE(6,104)
0007      103 FORMAT(1H-,36X,'NOX',11X,'NOX',
          &/13X,'VSC', 9X,'C',10X,'PPM DRY',4X,
          &'LB/10**6BTU',6X,'RUN #')
0008      104 FORMAT(1H1,25X,'SOUTHWESTERN PUBLIC SERVICE CO.',
          &/35X, 'SYSTEM LAB',/8X,
          &23X,'OXIDES OF NITROGEN')
          WRITE(6,103)
0009      2 CONTINUE
0010      TC = 0.
0011      TPPM = 0.
0012      TPDBTU = 0.
0013      DO 3 I=1,NSAMP
0014      READ (5,101) J,VF,PF,TF,PI,TI,AM,02
0015      101 FORMAT (3X,A4,4X,F5.0,5X,F5.2,5X,F4.0,5X,F5.2,
0016      &5X,F4.0,5X,F6.1,2X,F4.1)
          VSC=17.65*(VF-25.)*((PF/TF)-(PI/TI))
          C = 0.000062*AM/VSC
          PPM = C * 10.**9./(2.59*46.01)
          PDBTU = C * F * 20.9/(20.9 - 02)
          WRITE(6,102) VSC,C,PPM,PDBTU,J
0022      102 FORMAT(/10X,F8.2,2X,F10.7,4X,F8.2,6X,F8.6,8X,A5)
          TC = TC + C
          TPPM = TPPM + PPM
          TPDBTU = TPDBTU + PDBTU
0026      3 CONTINUE
          AC = TC / NSAMP
          APPM = TPPM / NSAMP
          APDBTU = TPDBTU / NSAMP
          WRITE(6,105)
0031      105 FORMAT(1H ,20X,'-----',
          &5X,' ',5X,'-----')
          WRITE(6,106)AC,APPM,APDBTU
0032      106 FORMAT(1H ,4X,'AVERAGE',8X,F10.7,4X,F8.2,6X,F8.6)
0033      READ(5,100)F,NSAMP
0034      IF(F.NE.0.)GO TO 2
0035      READ(5,108)PLANT(1),PLANT(2),PLANT(3),UNITNO,SAMLO(1),SAMLO(2),
0036      &SAMLO(3),SAMDAT(1),SAMDAT(2),
          &UTLOAD,SAMDAT(3),SAMDAT(4)
0037      108 FORMAT(5X,3A4,9X,A4,15X,3A4,
          &/12X,2A4,11X,A4,
          &/20X,2A4)
0038      WRITE(6,109)PLANT(1),PLANT(2),PLANT(3),PLANT(4),UNITNO,
          &SAMDAT(1),SAMDAT(2),
          &UTLOAD,SAMDAT(3),SAMDAT(4)
0039      109 FORMAT(///,5X,'PLANT',4A4,'STACK NO',A4,
          &/5X,'SAMPLE DATE ',2A4,' UNIT LOAD',A4,'MW',
          &/5X,'SAMPLE RECOVERY DATE ',2A4,
          &///31X,' ',
          &/36X,' ANALYZED BY')
          GO TO 30
0040      999 STOP
0041      END
0042

```

TABLE IV-II. PROGRAM FOR METHOD 7 CALCULATIONS

NO_x CALCULATIONS

$$V = (17.64 \frac{^{\circ}\text{R}}{\text{in. Hg}}) (V_f - 25 \text{ ml}) \left(\frac{P_f}{T_f} \frac{P_i}{T_i} \right) \quad (23)$$

where

- V_{sc} = Sample volume at standard conditions (dry basis), ml
- V_f = Volume of flask and valve, ml
- P_f = Final absolute pressure of flask, inches Hg
- P_i = Initial absolute pressure of flask, inches Hg
- T_f = Final absolute temperature of flask, $^{\circ}\text{R}$
- T_i = Initial absolute temperature of flask, $^{\circ}\text{R}$
- $17.64 = \frac{\text{Temperature standard} = (528 \text{ } ^{\circ}\text{R})}{\text{Pressure standard} = 29.92 \text{ in. Hg.}}$

$$C = (6.2 \times 10^{-5} \frac{\text{lb/scf}}{\text{ug/ml}}) \left(\frac{m}{V_{sc}} \right) \quad (24)$$

where

- C = Concentration of NO_x as NO₂ (dry basis) lb scf
- m = Mass of NO₂ in gas samples, ug
- V_{sc} = Sample volumes at standard conditions (dry basis), ml.

NO_x CALCULATIONS

To convert the concentration of NO_x in lb/dscf to lb/10⁶ Btu, the following formula from 60.45e (1) was used:

$$E = CF \frac{20.9}{20.9 - \%O_2} \quad (25)$$

where

- E = lb/10⁶ Btu
- C = lb/dscf
- F = F factor as calculated in 40 CFR 60.45f (5).

To convert C , the concentration of NO_x as NO₂ (dry basis, in lb/scf to parts per million), use the following:

$$\text{PPM} = C * 8.391662 * 10^6 \quad (26)$$

1. Reagents For NO_x Test:

NO_x Absorbing Solution: diluted 3 ml concentrated sulfuric acid to 1 liter with DI water. Mixed well. Added 3 ml 30 percent H₂O₂. Did not expose to heat or direct sunlight.

1 N Sodium Hydroxide: dissolved 42 grams NaOH in distilled water and diluted to 1 liter.

Phenoldisulfonic Acid: dissolved 25 grams pure white phenol in 150 ml concentrated sulfuric acid on a steam bath. Cooled, added 75 ml fuming sulfuric acid and heated at 100°C for two hours. Stored in a dark, glass-stoppered bottle.

Potassium Nitrate Standard Solution: dried potassium nitrate (KNO_3) at 105° to 110°C for two hours and cooled in a desiccating chamber. Dissolved 0.5495 grams KNO_3 in distilled water and diluted 1 liter in a volumetric flask. For a working standard, diluted 10 ml to 100 ml with water to obtain the working standard of 25 ug/ml (.025 mg/ml).

Using 10 ml of working standard yielded 250 ug NO_2 (.25 mg NO_2); other amounts equaled the following:

20 ml	yielded	500 ug NO_2	(.50 mg NO_2).
30 ml	yielded	750 ug NO_2	(.75 mg NO_2).
40 ml	yielded	1000 ug NO_2	
50 ml	yielded	1250 ug NO_2	
60 ml	yielded	1500 ug NO_2	
70 ml	yielded	1750 ug NO_2	
80 ml	yielded	2000 ug NO_2	
90 ml	yielded	2250 ug NO_2	
100 ml	yielded	2500 ug NO_2	
110 ml	yielded	2750 ug NO_2	
120 ml	yielded	3000 ug NO_2	

M. Calibrations and Standardizations

1. Flask Volume: The volume of the collection flask was determined prior to sampling. To do this, the flask and stopcock assembly were filled with water. Then the volume of water contained in the flask was measured and recorded to ± 10 ml.

2. Spectrophotometer: Calibration of the wavelength scale on the photometer was done under any of the following conditions:

If there was any replacement of parts such as lamps, etc., on the photometer.

If it had been six months since the last time it was checked.

The calibration procedure involved the drawing of a curve with transmittance versus wavelength. The System Lab maintained a set of solutions to be used for these wavelength checks. The instructions provided with the standard solutions were followed in drawing up these graphs. All of the data collected for wavelength checks and maintenance records were kept in an instrument quality control record book.

3. Quality Assurance: Three control charts were maintained to assure the quality of the data reported for NO_x determinations (see Figure IV-51). One control chart was set up for each of the three standard solutions used. Each chart plotted the results of the standard sample versus the date of analysis. The results of the analysis of the standard sample were plotted as percent difference from the true value.

QUALITY ASSURANCE

[illegible]

FIGURE IV-51. ANALYSIS OF STANDARDS FOR NO_x

The percent difference was calculated using the following formula:

$$d = \frac{C_m - C_t}{C_t} \times 100 \quad (27)$$

where

d = % difference

C_m = measured concentration

C_t = true concentration.

Comments were placed on the control chart when a new batch of standards was made up or a new reagent was used. Interpretation of the quality control charts is discussed under Quality Control and Assurance.

H. Quality Control and Assurance

This section deals with the mechanics of drawing up a quality control chart and how to use the chart to assure good quality data. The criteria for field equipment is also presented.

The performance criteria necessary for equipment used in sampling for particulate, SO_2 , and NO_x are listed in Tables IV-III, IV-IV, and IV-V. These criteria were adhered to in purchasing and calibrating equipment for the special test program, eliminating large undetected bias in sampling and analysis.

Control charts were used in SO_2 and NO_x analyses to verify and document the quality of the data collected. The percent difference from a standard value or an original value was plotted against time. The upper and lower control limits were either calculated using the standard deviation as a basis or were assigned a value from literature on the subject.

There were two types of control charts for SO_2 analysis. The first type involved the measurement of standard SO_2 samples. The value obtained on each standard was used to calculate the percent d ; this value was then plotted on separate charts for each standard. The second type of control chart was used to assure a consistent quality of barium perchlorate to titrate with. Before each set of analysis, the barium perchlorate normality was checked with standard sulfuric acid and then the percent d from the original value was calculated and plotted. Both of the control charts for SO_2 had 5 percent control limits until enough data was obtained to develop other limits. This limit is suggested in the "Guidelines For Development of a Quality Assurance Program: Volume V - Determination of Sulfur Dioxide Emissions From Stationary Sources" (3).

Table IV-III: Suggested Performance Criteria
For Particulate Sampling Equipment

Dry Gas Meter	Calibration factor between 0.98 and 1.02.
Thermometers and Temperature Gauges	Capable of measuring 1°C or 2°F intervals; must agree within 1.5% of the absolute temperature as measured by a standard in-glass mercury thermometer.
Stack Temperature Measuring System	Digital pyrometer with chromel-alumel thermocouple; must agree within 1.5% of the absolute temperature as measured by a standard in-glass mercury thermometer.
Sampling Train Leakage.	Less than 0.02 ft. ³ /minute at 15-inch Hg.
Sample Box Heater	Capable of maintaining a temperature of 250°F ± 25°F at laboratory conditions.
Probe Heating System	Capable of reaching and maintaining a temperature of 250°F ± 25°F at lab conditions inside the probe in approximately 30 minutes.
Magnehelic	Capable of reading within 5% of an incline gauge oil manometer.
Probe Nozzle Diameter	Range of three different diameter measurements less than 0.010 cm.
Analytical Balance	Weigh a standard weight within ± 0.3 mg. of its stated weight.

Table IV-IV. Suggested Performance Criteria
For SO₂ Sampling Equipment

Dry Gas Meter	Calibration factor between 0.98 and 1.02.
Thermometers and Temperature Gauges	Capable of measuring 1°C or 2°F intervals; must agree within 1.5% of the absolute temperature as measured by a standard in-glass mercury thermometer.
Stack Temperature Measuring System	Digital pyrometer with chromel-alumel thermocouple; must agree within 1.5% of the absolute temperature as measured by a standard in-glass mercury thermometer.
Sampling Train Leakage	1% of sampling flow rate at 10-inch Hg.
Probe Heating System	Capable of reaching and maintaining a temperature of 250°F ± 25°F at lab conditions inside the probe in approximately 30 minutes.

Table IV-V. Suggested Performance Criteria
For NO_x Sampling Equipment

Thermometers and Temperature Gauges	Capable of measuring 1°C or 2°F intervals; must agree within 1.5% of the absolute temperature as measured by a standard in-glass mercury thermometer.
U-Tube Manometer	36-inch mercury manometer with 0.1-inch divisions.
Flask Volume	Known within ± 10 ml.
Spectrophotometer	Wavelength should check out to within ± 5 nm.

For NO_x analysis, a control chart similar to the standard chart for SO_2 was used. The percent d was calculated and plotted against time. The control limits were arbitrarily set at 12 percent. After a sufficient number of analyses were run, the standard deviation was calculated, then the control limit was set at three times the standard deviation.

Drawing and maintaining control charts is not good enough to assure high quality data. Action must be taken to correct situations when the procedure is out of control. The following list indicates the criteria SPS used to recognize a process out of control.

One point fell outside the upper or lower control limit.

Two consecutive points fell between the warning and control limits.

Three consecutive points fell outside the standard deviation range.

The System Lab also participated in the Stationary Source Quality Assurance Survey conducted by the EPA. The results of these standard analysis are kept on file as a part of the quality assurance program for SO_2 , NO_x , and particulate analysis. The form used by the System Lab for the NO_x particulate survey is included here (Figure IV-52).

Calibrated By _____

Dry Gas Meter Correction Factor _____

in H₂O (ΔH) DG_i DG_f T_{in i} T_{out i} T_{in f} T_{out f}

(ft)³ (ft)³ (°F) (°F) (°F) (°F)

Amb. Temp. Initial Amb. Temp. Final Vac. Time
minutes minutes Pbar Run No.

Dry Gas Meter Correction Factor

Calibrated By

Barometric Pressure

in H_2O (ΔH)
$$DG_i^3 \text{ (ft}^3\text{)}$$
$$DG_f \quad (\text{ft}^3)$$
 $T_{in\ i}$
(°F)
$$T_{out,i} \quad (^{\circ}F)$$
 T_{inf}
(°F)
$$T_{out\ f} \text{ (}^{\circ}\text{F)}$$

Amb.
Temp.
Initial

[illegible]

Vac. Hq.

Time
minute

Run No.

[illegible]

FIGURE IV-52. METHOD 5 DRY GAS METER INTER-LABORATORY SURVEY

V. SPS TEST PHASES AND RESULTS

A. Phase I

During the week of December 11-15, 1978, SPS performed the first series of tests to measure mass emissions of particulate, sulfur dioxides, and oxides of nitrogen. The original work plan called for simultaneous sampling at five locations; however, procurement problems with equipment prohibited completion of the outlet sampling trains. For particulate and SO_2 samples it was believed the test would result in better quality data if only three stations (both inlets and the stack) were sampled. The sampling procedure for oxides of nitrogen required less manpower and equipment than sampling for SO_2 and particulate; therefore, sampling for NO_x was performed at all five locations (two inlet, two outlet, one stack).^x

A crew of test personnel from SPS power plants was assembled at Harrington Station for the first week-long series of tests. Approximately 26 stack sampling team members participated in the program. SPS organized a stack sampling manual for personnel assigned to the special testing phases. By taking samples at three locations rather than five, experienced personnel were able to work with the less experienced ones. As a result, personnel at all the sampling locations had some degree of experience during the remaining tests.

The results of the first inlet testing compare favorably with the theoretical inlet grain loading (the theoretical inlet grain loading is estimated and based upon generation of 70 percent fly ash). The percent fly ash has never been accurately determined but has been estimated to be between 70 and 80 percent.

During the Phase I test, the heater on the Inconel probe being used at the stack location failed. It was later determined that condensation forming in the probe contributed additional particulates to the probewash; this solution was greenish in color. It was later analyzed and determined to be high in nickel. The conclusion was that the acid moisture reacted with the probe and thereby biased the test result. These conditions were corrected for the second series of tests. The results of the NO_x tests were very consistent across the baghouse. Results of the sulfur dioxide tests on the stack compare favorably with the stoichiometric calculation for sulfur dioxide. SO_2 measurements at inlet locations were erratic due to high negative duct pressure.

Analyses of the Phase I special tests are shown in Table V-1. Results confirmed that Unit 2 can operate in compliance with the existing New Source Performance Standard of $.1 \text{ lb}/10^6 \text{ Btu}$. However, problems in the testing with the unheated probe caused results to be biased (high). Concentrations of NO_x were determined to be consistent at all five sampling locations. As a result, SPS and EPA decided the testing program could be better managed by excluding NO_x from future testing. Field notes are included in Appendix C.

B. Phase II

The second series of special tests to be performed by SPS was initially scheduled for May, 1979. Due to a decision to rebag the fabric filter system, special testing was rescheduled for October 8-12, 1979.

In September of 1978, the Harrington Station baghouse began to experience a higher than normal failure of bags. Examination of the problem indicated that two items needed immediate attention: (1) the

Table V-1. Phase I Test Results

Results of Particulate Testing

Run #	East Inlet gr/SCF	West Inlet gr/SCF	Theoretical Inlet+ gr/SCF	Stack*** gr/SCF	lb/10 ⁶ Btu
1*	2.28	2.74	2.09	.053	.106
2*	2.04	2.26	1.96	.050	.097
3**	1.67	1.63	2.27	.033	.061

*sootblowing continuously

**not sootblowing

***The concentrations of particulate obtained from the stack are biased high because a reaction took place in the unheated Inconel probe liner.

+assumes 70% fly ash, 30% bottom ash and economizer ash; no consideration for sootblowing.

Results of NO_x Testing

Run #	East Inlet Method 7	East Outlet Method 7	West Inlet Method 7	West Outlet Method 7	Stack Method 7
1	.68	.64	.61	.62	.63
2	.71	.68	.59	.62	.66
3	.67	.71	.62	.62	.64

Results of SO₂ Testing

Run #	East Inlet* Method 6 lb/10 ⁶ Btu	West Inlet* Method 6 lb/10 ⁶ Btu	Stoichiometric lb/10 ⁶ Btu	Stack Method 6 lb/10 ⁶ Btu
1	.53	.36	.76	.73
2	.59	.32	.84	.78
3	.62	.20	.88	.84

*These concentrations are suspected of being low because of the high negative pressure pulling the absorbing solutions forward, thereby resulting in the absorbed SO₂ not being analyzed.

**Assumes all sulfur is converted to SO₂.

control of the deflation pressure during the cleaning cycle and (2) tension on the bag. By October, a program had begun to redesign the deflation pressure control system and bags were re-tensioned.

SPS then became concerned about the accelerated rate of bag failures occurring in the Unit 2 baghouse. These failures were of two types; small pinhole-type failures near the bottom cuff, and failures described as "blowouts" where a whole section of the bag would burst and fray. Based on information derived from an in-house fabric assessment program and an EPA mobile baghouse study, SPS decided to rebag the entire baghouse with new fabrics before the 1979 summer peak. The west baghouse was rebagged June 16, 1979 and the east on June 30, 1979. Figure V-1 shows the types of fabrics used to rebag the baghouse.

Another adjustment that resulted from the EPA mobile baghouse study was resetting the cleaning cycle and increasing the frequency of shake on two compartments. The adjustment in frequency of shake was accomplished by changing the size of the shaker's drive mechanism pulley.

A relative flow comparison between compartments with a standard shake and an accelerated shake indicated much improvement in cleaning and pressure drop. Larger pulleys were placed in additional compartments and two problems resulted: (1) the limitation of shaker motor horsepower could not be exceeded and (2) the shaker pillow block system began to fail. It was determined that a 5.3-inch pulley in place of the 4.0-inch pulley provided enough frequency increase and was within horsepower limitations. The pillow block assemblies were redesigned and replaced before the second special test; no recent problems have been noted.

With these changes having been completed, SPS conducted the Phase II special testing in October, 1979. The experience gained in the first series of tests enabled SPS personnel to conduct tests at all five sample locations as originally specified by the EPA. Results of the Phase II testing are shown in Table V-II.

Data from the outlet sample locations were not of as good a quality as data from stack sample locations. This difference may be due to a poor outlet sampling location and an extra long port extension. It was very difficult to sample these locations without picking up material in the probe nozzle, and a very small quantity can significantly affect the results.

High negative pressure still affected the Phase II SO₂ measurements, although results were much improved over the first test. This problem was attributed to negative pressure pulling the absorbing solution forward in the sampling train.

C. Phase III

Southwestern's third special test was performed the week of March 24-27, 1980. Twenty SPS employees participated in the week-long test for SO₂ and particulate. A number of difficulties were encountered, causing several delays; however, it was felt that the testing was concluded in a successful manner.

Adjustments made to the baghouse prior to the Phase III testing included the following:

All of the west baghouse compartments were retensioned after several of them failed to shake. This problem was attributed to over-

WEST BAGHOUSE		EAST BAGHOUSE	
1 *	2 *	15 **	16 **
3 *	4 *	17 **	18 **
5 *	6 Menardi-Southern Teflon Test Bags	19 Original bags equipped with special shaker mechanism	20 Fabric Filters All-filament Teflon
7 1 Nomex All-Spun; 2 Nomex Comb. 5 Crisoflex 446; 4 Crisoflex 449 ***	8 *	21 Criswell 442 Teflon B Test Bags	22 34 Acid Flex; 34 Tri-Treat Balance: Original Bags
9 * (Warp In)	10 *	23 Fabric Filters 502 Tri-Treat Test Bags	24 Globe-Albany Nomex
11 *	12 *	25 Criswell 449 Teflon B Test Bags	26 **
13 *	14 *	27 **	28 **

*Criswell 442 Teflon B, 0.30kg (10.5 oz.) (rebagging complete).
 **Criswell 449 Tri-Treat: 0.40kg (14 oz.) (rebagging complete).

FIGURE V-1. FABRIC INSTALLATION, JULY, 1979

Table V-II. Phase II Test Results

Results of Particulate Testing:

Run	East		West		Theo-		East		West	
	Inlet	gr/scf	Inlet	gr/scf	retical	gr/scf [±]	Outlet	gr/scf	Outlet	gr/scf
No.										
1	1.40		1.41		1.64		.014		.012	
2	1.61		1.41		1.64		.014		.024	
3	1.61		.74*		1.69		.027		.028	
									.011	
									.010	
									.020	
									.021	
									.022	

Results of SO₂ Testing:

Run	East		West		Theo-		East		West	
	Inlet	lb/10 ⁶ Btu	Inlet	lb/10 ⁶ Btu	retical	lb/10 ⁶ Btu	Outlet	lb/10 ⁶ Btu	Outlet	lb/10 ⁶ Btu
No.										
1	.71		.74		1.03		.70		.76	
2	.74		.79**		.95		.79		.82	
3	.66		.71**		.99		.78		.78**	
									.79	
									1.01	
									.96	

[±] Based upon stoichiometric flue gas flow, coal flow, ash analysis and 70% fly ash generation and 30% bottom ash and economizer ash generation.

* One sample of probe wash was lost during transport from sample site to the lab, causing this sample to be biased low.

** Based upon only one of two samples for this run.

tightening of turnbuckles attached to the shaker tube support member. These turnbuckles were installed during the October, 1979 annual over-haul to correct sagging problems. Apparently, they were over-tightened, which in turn over-tensioned the bags, resulting in an overload condition on the shaker motor.

During the month of March, three outlet damper drives had to be repaired. As in all previous outlet damper drive failures the brass nut had failed. In order to lengthen damper drive life, tests were performed to see if the damper drive stroke could be shortened without affecting the pressure drop across the outlet damper.

Results of the Phase III testing are shown in Table V-III. Several problems were encountered during the testing. Inlet testing teams had considerable problems with the fritted supporting disks breaking, causing the filters to tear. This accounts for the low grain loading experienced on the various runs. The last run at the stack location failed the leak test at the end of the run. SPS determined that a connector glassware clamping the hot box lost tension and leaked. A review of the raw data indicates this probably happened at the first point in the last port.

A review of the SO_2 results indicates that problems still existed with sampling the highly negative pressure ducts with the SO_2 impingers. Improvements in testing procedures had been made by the third test, but some problems still remained to be solved. These are itemized in Appendix C.

D. Phase IV

Maintenance and operation changes between the third and fourth special tests ranged from cleaning boiler slag out of the Unit 2 economizer and air preheaters to repairs to shaker tube flange bearings. The bags in compartment 23 were changed out during April, due to multiple pinhole failures in the upper portion of the bags. In addition, seven shaker mechanisms were repaired in June.

During the Phase IV testing period in August, 1980, the baghouse encountered no major upsets or operating problems. All of the shakers were in service and operating in a normal fashion. The temperatures, flows and pressure drops encountered were all within normal operating limits.

The opacity, however, was higher than normal, being approximately 8 percent compared to 5 percent previously. (It is felt that this rise was due to bag bleeding in several compartments of test fabrics).

During the test period erratic opacity spikes were observed that seemed to correspond with the cleaning cycle of compartment 27. The compartment was left in service throughout the test awaiting more definite indications of failure since the opacity spikes did not follow the clean cut blowout pattern previously observed. On August 29, 1980 the compartment was removed from service and four blown bags and two other damaged bags were replaced.

Problems with sampling SO_2 at inlets were corrected and results were comparable to stack SO_2 (see Table V-IV).

E. SPS/GCA Test

A joint test was conducted by SPS and its subcontractor, GCA Corporation, on June 4, 5, and 6, 1980. SPS ran simultaneous tests for particulate and GCA tested for SO_3 and particle sizing.

Table V-III. Phase III Test Results

Results of Particulate Testing:						
Run No.	East		West		East	
	Inlet	gr/scf	Inlet	gr/scf	Theoretical	Outlet
					gr/scf**	gr/scf
						Stack
						lb/10 ⁶ Btu
1	*	.	*	.	1.82	.008
2	1.43		1.35		1.95	.013
3	.97		1.14		2.03	.011
4	*	.	1.50		2.04	.007
						.012
						.038
						.022
						.020
						.009
						.017±
						.024
						.032
						.044±±

Results of SO ₂ Testing:						
Run No.	East		West		East	
	Inlet	lb/10 ⁶ Btu	Inlet	lb/10 ⁶ Btu	Theoretical	Outlet
					lb/10 ⁶ Btu	lb/10 ⁶ Btu
						Stack
						lb/10 ⁶ Btu
1	.76		.69		.82	.70
2	*		.67		.81	.69
3	*		.67		.81	.65
						.68
						*
						*
						.81
						.75
						.78

* No data

** Based upon stoichiometric flue gas flow, coal flow, ash analysis and 70% fly ash generation and 30% bottom ash and economizer ash generation; (dry basis).

± Not isokinetic

±± Failed leak check test

Table V-IV. Phase IV Test Results

Results of Particulate Test:

Run	East Inlet	West Inlet	Theoretical	Stack	
No.	gr/scf	gr/scf	gr/scf**	gr/scf	lb/10 ⁶ Btu
1	1.64	1.92	1.99	.016	.031
2	1.88	1.67	1.97	.009	.018
3	1.35	1.44	1.96	.015	.030

Results of SO₂ Test:

Run	East Inlet	West Inlet	Theoretical	Stack
No.	lb/10 ⁶ Btu	lb/10 ⁶ Btu	lb/10 ⁶ Btu	lb/10 ⁶ Btu
1	.84	.76	.96	.84
2	.89	.79	.94	.95
3	.88	.78	.96	.83

** Based upon stoichiometric flue gas flow, coal flow, coal analysis and 70% fly ash generation (dry basis).

SPS testing began at 10:50 a.m. on June 4, 1980. No significant problems occurred during the testing. The east inlet probe liner slipped as the probe was being withdrawn from the duct. This caused the train to fail the leak check; however, it is not believed that results were significantly affected. The test concluded at about 12:30 p.m.

GCA began the impactor testing at the stack location at about 7:00 a.m. Goksoyr-Ross was run for 1 hour. No condensation occurred in the condenser coil indicating the low concentration of SO_3 . A discussion with GCA indicated that a much longer run on the stack might give better resolution of SO_3 . This was not predicted on the inlets because of the high grain loading of particulate. GCA also began the inlet impactor runs about 12:30 p.m. and expected to run until about 4:00 p.m.

No particular problems were experienced by SPS. The stack run blew two fuses while in the fourth port, which was noted on a data sheet and sampling time and volume were kept up. In order to collect a larger particulate sample, a larger nozzle was selected for the third run. This allowed a larger sample and minimized problems associated with verifying a small particulate sample. The baghouse experienced failures in the shaker mechanism.

The testing during these two tests were considered by SPS the most successful of any conducted during the program. The inlet particulate concentrations were not as consistent as desired which can be attributed to the lack of idealness in the sampling location. The inlet SO_2 emissions were consistent and compared very well with the stack results. Much difficulty was experienced with the highly negative duct pressure in previous tests. Results of the SPS/GCA test are shown in Table V-V.

Table V-V. Results of SPS/GCA Joint Test

Results of Particulate Test:

Run No.	East	West	Theoretical gr/scf**	Stack	
	Inlet gr/scf	Inlet gr/scf		gr/scf	lb/10 ⁶ BTU
1	*	2.30	2.13	.015	.032
2	1.92	1.86	2.22	.019	.037
3	1.38	1.31	1.95	.008	.016

*Invalid test

**Based upon stoichiometric flue gas flow, coal flow, coal analysis and 70% fly ash generation (dry basis).

VIII. REFERENCES

1. Ladd, K. L., Faulkner, G. R. and Kunka, S. L., "Fabric Filter System Study: First Annual Report", EPA-600/7-79-183, August 1979.
2. Ladd, K. L., Chambers, R. C., Plunk, O. C., and Kunka, S. L., "Fabric Filter System Study: Second Annual Report", prepared under EPA Contract No. 68-02-2659, August 1980.
3. "Guidelines For Development of a Quality Assurance Program: Volume V - Determination of Sulfur Dioxide Emissions From Stationary Sources."

Appendix A
Coal Analysis

COMMERCIAL TESTING & ENGINEERING CO.

GENERAL OFFICES: 228 NORTH LA SALLE STREET, CHICAGO, ILLINOIS 60601 AREA CODE 312 726-8434

WESTERN DIVISION MANAGER
LLOYD W. TAYLOR, JR.



PLEASE ADDRESS ALL CORRESPONDENCE TO:
10775 EAST 51st AVE., DENVER, COLO. 80239
OFFICE TEL. (303) 373-4772

► SOUTHWESTERN PUBLIC SERVICE COMPANY
P.O. Box 1261
Amarillo, Texas 79170

January 12, 1979

Sample Identification
by

Kind of sample reported to us	Coal	Southwestern Public Service Co.
Sample taken at	Harrington Station	Sample No. 30
Sample taken by	Southwestern Public Service Co.	Feeder A - E
Date sampled	12-14-78	Air Dried 18 hours
Date received	1-4-79	% ADL = 18.00
		Charge Account #606-1501.0402
		P. O. No. 766-060F

Analysis report no. 72-80117

PROXIMATE ANALYSIS

	As Received	Dry Basis
% Moisture	29.20	xxxx
% Ash	5.62	7.94
% Volatile	31.63	44.68
% Fixed Carbon	33.55	47.38
	100.00	100.00
Btu	8437	11916
% Sulfur	0.30	0.42

ULTIMATE ANALYSIS

	As Received	Dry Basis
% Moisture	29.20	xxxx
% Carbon	49.03	69.25
% Hydrogen	3.45	4.87
% Nitrogen	0.76	1.07
% Chlorine	0.01	0.01
% Sulfur	0.30	0.42
% Ash	5.62	7.94
% Oxygen (diff)	11.63	16.44
	100.00	100.00

cc: Ken Lindemann
Jerry Hermesmyer
Royce Kelly
Steve Jones
Charles Bagwell
Morris Rantz
Olton Plunk ✓

REPRESENTATIVE COAL ANALYSIS PHASE I

Respectfully submitted,
COMMERCIAL TESTING & ENGINEERING CO.

G. D. PALMER, Manager, Denver Laboratory



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COMMERCIAL TESTING & ENGINEERING CO.

GENERAL OFFICES: 228 NORTH LA SALLE STREET, CHICAGO, ILLINOIS 60601 AREA CODE 312 725-8434

WESTERN DIVISION MANAGER
LLOYD W. TAYLOR, JR.



PLEASE ADDRESS ALL CORRESPONDENCE TO:
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OFFICE TEL. (303) 373-4772

▷ SOUTHWESTERN PUBLIC SERVICE COMPANY
P.O. Box 1261
Amarillo, Texas 79170

November 27, 1979

Sample identification
by

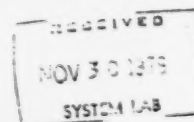
Kind of sample reported to us	Coal	Southwestern Public Service Co.
Sample taken at	xxxxxx	Sample No. 45
Sample taken by	Southwestern Public Service Co.	% Air Dry Loss = 10.38
Date sampled	10-11-79	Composite Baghouse
Date received	11-1-79	Charge Account # 606-1501.0402
		P.O. No. 15326-75

Analysis report no. 72-88356

PROXIMATE ANALYSIS			ULTIMATE ANALYSIS		
	As Received	Dry Basis		As Received	Dry Basis
% Moisture	26.91	xxxx	% Moisture	26.91	xxxx
% Ash	4.69	6.42	% Carbon	51.32	70.21
% Volatile	32.44	44.38	% Hydrogen	3.71	5.08
% Fixed Carbon	35.96	49.20	% Nitrogen	0.65	0.89
	100.00	100.00	% Chlorine	0.01	0.01
			% Sulfur	0.36	0.49
Stu	8850	12108	% Ash	4.69	6.42
% Sulfur	0.36	0.49	% Oxygen (diff)	12.35	16.90
				100.00	100.00

cc: Steve Jones
Ken Lindemann
Olton Plunk
Ronnie Blythe
Steve Thurman

REPRESENTATIVE COAL ANALYSIS PHASE II



Respectfully submitted,
COMMERCIAL TESTING & ENGINEERING CO.

G. D. Palmer

G. D. PALMER, Manager, Denver Laboratory



Charter Member

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COMMERCIAL TESTING & ENGINEERING CO.

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WESTERN DIVISION MANAGER
LLOYD W. TAYLOR, JR.



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OFFICE TEL. (303) 373-4772

► SOUTHWESTERN PUBLIC SERVICE COMPANY
P.O. Box 1261
Amarillo, Texas 79170

April 25, 1980

Sample identification
by

Kind of sample reported to us Coal
Sample taken at xxxxx
Sample taken by Southwestern Public Service Company
Date sampled 3-27-80 (2,3,4,5 p.m.)
Date received 4-15-80

Southwestern Public Service Co.

Sample No. 71
% ADL = 10.37

Analysis report no. 72-92452

PROXIMATE ANALYSIS

	As Received	Dry Basis
% Moisture	27.27	xxxx
% Ash	4.98	6.85
% Volatile	32.46	44.63
% Fixed Carbon	35.29	48.52
	100.00	100.00
Btu	8840	12155
% Sulfur	0.35	0.48

ULTIMATE ANALYSIS

	As Received	Dry Basis
% Moisture	27.27	xxxx
% Carbon	51.45	70.74
% Hydrogen	3.42	4.70
% Nitrogen	1.04	1.43
% Chlorine	0.01	0.01
% Sulfur	0.35	0.48
% Ash	4.98	6.85
% Oxygen (diff)	11.48	15.79
	100.00	100.00

cc: Ken Lindemann
Ronald Blythe
Steve Jones
Olton Plunk
Leo Fajardo

REPRESENTATIVE COAL ANALYSIS PHASE III

RECEIVED

APR 30 1980

SYSTEM LAB

Respectfully submitted,
COMMERCIAL TESTING & ENGINEERING CO.

G. D. Palmer
G. D. PALMER, Manager, Denver Laboratory



GDP/md/jh
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MOBILE, AL - NEW BETHLEHEM, PA - NEW ORLEANS, LA - NORFOLK, VA - FALLS CHURCH, VA - PITTSBURGH, PA - RICHMOND, VA - SALINA, UT - ST. LOUIS, MO - TAMPA, FL - TOLSON, OH - VANCOUVER, B.C. CAN.

COMMERCIAL TESTING & ENGINEERING CO.

GENERAL OFFICES: 228 NORTH LA SALLE STREET, CHICAGO, ILLINOIS 60601

AREA CODE 312 726-8434

WESTERN DIVISION MANAGER

GAIL D. PALMER


 PLEASE ADDRESS ALL CORRESPONDENCE TO:
 10775 EAST 51st AVE., DENVER, COLO. 80239
 OFFICE TEL: (303) 373-4772

SOUTHWESTERN PUBLIC SERVICE COMPANY

September 8, 1980

P.O. Box 1261

Amarillo, Texas 79170

Sample identification
by

Kind of sample reported to us Coal

Sample taken at xxxxx

Sample taken by Southwestern Public Service Co.

Date sampled 8-26-80

Date received 9-2-80

Southwestern Public Service Co.

Sample No. 3

Time: 3:20 pm & 4:30 pm

Moisture Loss = 17.41%

P. O. No. 15975-673

Analysis report no. 72-97065

PROXIMATE ANALYSIS

	As Received	Dry Basis
% Moisture	27.64	xxxx
% Ash	4.94	6.83
% Volatile	32.00	44.23
% Fixed Carbon	35.42	48.94
	100.00	100.00
Btu	8763	12110
% Sulfur	0.42	0.58

ULTIMATE ANALYSIS

	As Received	Dry Basis
% Moisture	27.64	xxxx
% Carbon	50.50	69.79
% Hydrogen	3.60	4.98
% Nitrogen	0.72	1.00
% Chlorine	0.01	0.02
% Sulfur	0.42	0.58
% Ash	4.94	6.83
% Oxygen (diff)	12.17	16.80
	100.00	100.00

cc: Ken Lindemann

Ronnie Blythe

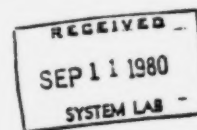
Steve Jones

Dean Metcalf

Olon Plunk

REPRESENTATIVE COAL ANALYSIS

PHASE IV



GDP/md/pm

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 Respectfully submitted,
 COMMERCIAL TESTING & ENGINEERING CO.

G. D. PALMER, Manager, Denver Laboratory



Charter Member

 BILLINGS, MT. - BIRMINGHAM, AL. - CHARLESTON, WV. - CLARKSBURG, WV. - CLEVELAND, OH. - CONNEAUT, OH. - DENVER, CO. - GOLDEN, CO. - HELPER, UT. - HENDERSON, KY. - HOUSTON, TX. - JASPER, AL.
 KNOXSBORO, KY. - KNOX, TN. - NEW BETHLEHEM, PA. - NEW ORLEANS, LA. - NORFOLK, VA. - PALISADE, CO. - PINEVILLE, KY. - SALINA, UT. - SD. - HOLLAND, IL. - TOLEDO, OH. - VANCOUVER, B.C. CAN.

COMMERCIAL TESTING & ENGINEERING CO.

GENERAL OFFICES: 228 NORTH LA SALLE STREET, CHICAGO, ILLINOIS 60601

AREA CODE 312 728-8434

WESTERN DIVISION MANAGER
LLOYD W. TAYLOR, JR.



PLEASE ADDRESS ALL CORRESPONDENCE TO:
10775 EAST 51st AVE., DENVER, COLO. 80239
OFFICE TEL. (303) 373-4772

► SOUTHWESTERN PUBLIC SERVICE COMPANY
P.O. Box 1261
Amarillo, Texas 79170

July 21, 1980

Sample identification
by

Kind of sample reported to us	Coal	Southwestern Public Service Company
Sample taken at	XXXXXX	Composite
Sample taken by	Southwestern Public Service Company	Baghouse - Sample 75
Date sampled	6-5-80	P. O. No. 15863-75
Date received	7-8-80	% ADL = 12.57 (Reported by SPS CO.)

Analysis report no. 72-94929

PROXIMATE ANALYSIS

	<u>As Received</u>	<u>Dry Basis</u>
% Moisture	26.95	xxxx
% Ash	5.30	7.25
% Volatile	31.52	43.16
% Fixed Carbon	35.22	49.59
	<u>100.00</u>	<u>100.00</u>
Btu	3774	12012
% Sulfur	0.42	0.57

ULTIMATE ANALYSIS

	<u>As Received</u>	<u>Dry Basis</u>
% Moisture	26.95	xxxx
% Carbon	50.37	69.96
% Hydrogen	3.70	5.07
% Nitrogen	0.77	1.06
% Chlorine	0.00	0.00
% Sulfur	0.42	0.57
% Ash	5.30	7.25
% Oxygen (diff)	12.48	17.09
	<u>100.00</u>	<u>100.00</u>

cc: Ken Lindemann
Ronald Glythe
Environmental Group
Steve Jones
Joe Don Parrott

REPRESENTATIVE COAL ANALYSIS
SPS/GCA

Respectfully submitted,
COMMERCIAL TESTING & ENGINEERING CO.

G. D. Palmer

G. D. PALMER, Manager, Denver Laboratory



Charter Member

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Appendix B
Raw Field Data

	1	2	3	4	5	6	7	8	9	10	11	12	13	14
1 INLET PART LOADING EAST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
2 INLET PART LOADING EAST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
3 INLET PART LOADING WEST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
4 INLET PART LOADING WEST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
5 INLET PART LOADING EAST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
6 INLET PART LOADING EAST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
7 OUTLET PART LOADING WEST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
8 OUTLET PART LOADING WEST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
9 PART REMOVAL OFF EAST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
10 PART REMOVAL OFF WEST	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
11 INLET 502 EAST 1 HR AVG	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
12 INLET 502 EAST 1 HR AVG	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
13 INLET 502 WEST 1 HR AVG	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00
14 INLET 502 WEST 1 HR AVG	1.00	2.00	3.00	4.00	5.00	6.00	7.00	8.00	9.00	10.00	11.00	12.00	13.00	14.00

PHASE I

SOUTHWESTERN PUBLIC SERVICE CO.
PARTICULATE TEST SUMMARY

TEST NUMBER	1	2	3
ORSAT GAS ANALYSIS			
CO ₂	13.30	13.90	13.60
O ₂	0.0	0.0	0.0
N ₂	80.60	80.95	81.45
EXCESS AIR	37.24	32.16	29.90
DRY GAS M/L WT	30.16	30.45	30.37
PARTICULATE DATA			
BAROMETRIC PRESSURE	26.80	26.40	26.20
STATIC PRESSURE	-16.00	-16.40	-16.30
NOZZLE AREA	.30784E-03	.30784E-03	.30784E-03
PITOT TUBE COEFFICIENT	0.851	0.851	0.851
DISCHG. COEFF. AREA	0.998	0.998	0.998
TOTAL SAMPLING TIME	96.0	98.0	98.0
GAS SAMPLE VOL. METER COND.	77.61	80.71	83.67
MOISTURE COLLECTED	201.85	177.40	183.05
SAMPLE WEIGHT	12.3279	10.3713	7.5734
AVG STACK TEMP	799.8	793.0	805.3
AVG GAS METER TEMP	531.3	532.1	524.3
AVG SMOOT DELTA P	2.02	2.21	2.32
AVG SMOOT AREA	359.5	350.2	358.2
AVG LOAD	205.5	205.7	204.2
AVG COAL FLOW RATE	8525.	8437.	8517.
AVG BTU COAL	3503.1	3471.8	3479.2
AVG HEAT RATE	3.3	2.8	2.8
AVG OPACITY			

PARTICULATE RESULTS

SAMPLE VOL. DRY STD COND	69.340	70.937	71.631
PERCENT MOISTURE	12.09	10.56	10.77
NET STACK GAS MOL WT	29.87	29.13	29.04
ABSOLUTE DUCT PRESS	25.62	25.19	25.00
AVG DUCT VELOCITY	0.459	0.455	0.454
DUCT FLOW RATE STD COND	0.915	0.908	0.929
DUCT FLOW NET STACK COND	2.7433	2.2559	1.6114
PARTICULATE CONC	9050.19	7196.19	5075.92
PART EMISS RATE	2.5835	2.0727	1.4599
PART EMISS RATE	98.22	98.34	99.58
ISOKINETIC VARIATION			

B-2

SOUTHWESTERN PUBLIC SERVICE CO.
PARTICULATE TEST SUMMARY

TEST NUMBER

4

5

6

ORSAT GAS ANALYSIS

O₂ 12.45 13.30
CO 0.0 0.0
CO₂ 8.25 8.00
N₂ 81.30 81.50
EXCESS AIR 39.91 42.68
DRY GAS MUL WT 30.24 30.37

PARTICULATE DATA

BAROMETRIC PRESSURE 26.80 26.40 26.20
STATIC PRESSURE -19.80 -19.80 -19.80
NOZZLE AREA 0.379586-03 0.379586-03 0.379586-03
PITOT TUBE COEFFICIENT 1.003 1.003 1.003
DUCT CROSS SECT AREA 246.0 246.0 246.0
TOTAL SAMPLING TIME 98. 98. 98.
GAS SAMPLE VOL METED CUND 85.70 90.55 84.35
MOISTURE COLLECTED 18.50 210.10 20.15
SAMPLE WEIGHT 11.6257 100.31 7.1665
AVERAGE TEMPERATURE 52.5 55.3 54.9
AVG GAS METER TEMP 52.5 55.3 54.9
AVG 50 WHIT DELTA P 1.98 2.08 2.15
AVG LUAL 359.5 359.2 358.2
AVG CUAL FLUX RATE 295.5 295.7 297.2
AVG HU LUAL 3525. 3417. 3479.2
AVG HUAT RATE 3503.1 3417. 3479.2
AVG UPACITY 3.3 2.8 2.8

PARTICULATE RESULTS

SAMPLE VOL DRY STD CUND 76.727 79.473 72.466
PERCENT MOISTURE VOL X 10.15 11.48 12.59
WET STACK GAS MUL WT 25.04 25.28 25.93
ABSOLUTE DUCT PRESS IN HG 6.73 6.11 6.03
AVG DUCT FLOW DRY STD CUND 1046 DSCFM 0.479 0.465 0.463
DUCT FLOW WET STACK CUND 0.955 0.946 0.975
PARTICULATE CONC 2.3380 2.0432 1.6689
PART MISS RATE 8343.99 7081.91 5505.52
PART MISS RATE LB/1000 BTU 2.3619 1.921 1.5616
ISOKINETIC VARIATION 106.62 113.16 103.51

SOUTHWESTERN PUBLIC SERVICE CO.
PARTICULATE TEST SUMMARY

TEST NUMBER	7	8	9
OXSAT GAS ANALYSIS			
O2	13.40	14.41	13.56
CO	0.00	0.00	0.00
CO2	6.20	5.81	5.40
N2	80.40	79.77	81.03
EXCESS AIR	41.26	39.15	33.76
DRY GAS MOL WT	30.39	30.54	30.39
PARTICULATE DATA			
BAROMETRIC PRESSURE	26.70	26.29	26.06
NOZZLE AREA	0.90	1.00	0.89
PITOT TUBE COEFFICIENT	0.794	0.794	0.794
DUCT CROSS SECT AREA	1.006	1.006	1.006
TOTAL SAMPLING TIME	33.98	33.98	33.98
WET FLOW WET STACK COND	39.96	39.96	39.96
SAMPLE WEIGHT	76.15	70.10	80.50
AVG STACK TEMP	0.1267	0.1190	0.0764
AVG GAS METER TEMP	61.9	61.0	820.0
AVG SQ ROOT DELTA P	50.8	51.5	51.3
AVG DELTA H	1.19	1.18	1.20
AVG LOAD FLOW RATE	1.08	1.08	1.07
AVG DUCT FLOW RATE	35.2	35.2	35.2
AVG DUCT FLOW RATE	8525	8437	8517
AVG HEAT RATE	3503.1	3671.8	3472.2
AVG OPACITY	3.3	2.8	2.8
PARTICULATE RESULTS			
SAMPLE VOL DRY STD COND	36.471	36.508	35.958
WET STACK GAS MOL WT	29.29	29.39	29.20
ABSOLUTE DUCT PRESS	26.63	26.22	25.99
DUCT FLOW WET STD COND	0.888	0.871	0.870
DUCT FLOW WET STACK COND	1.090	1.079	1.070
PARTICULATE CONC	0.0531	0.0503	0.0528
PART EMISS RATE	370.1	350.7	361.1
PART EMISS RATE	98.09	99.53	98.12
ISOKINETIC VARIATION	LU/10**6 RTU	99.53	98.12

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 1														
10/11/79 0:30:0														
1 INTG GROSS GENERATION														
2 TOTAL INTG COAL FLOW														
3 IGNITER GAS FLOW-INTG														
4 MAIN GAS FLOW-INTG														
5 INLET TEMP EAST 1 HR AVG														
6 INLET TEMP WEST 1 HR AVG														
7 OUTLET TEMP EAST 1 HR AVG														
8 OUTLET TEMP WEST 1 HR AVG														
9 TEMP DIFF EAST 1 HR AVG														
10 TEMP DIFF WEST 1 HR AVG														
11 PRESS DIFF EAST 1 HR AVG														
12 PRESS DIFF WEST 1 HR AVG														
13 AIR FLOW EAST 1 HR AVG														
14 AIR FLOW WEST 1 HR AVG														
15														
16														
17														
18														
1	2	3	4	5	6	7	8	9	10	11	12	13	14	
MWH	TONS	MCF	MCF	DEG F	DEG F	DEG F	DEG F	DEG F	DEG F	IN H2O	IN H2O	MSCFM	MSCFM	
1:00 334.0	168.3	0.0	0.0	356.3	351.8	354.4	346.8	1.9	5.1	7.0	7.4	426.5	408.6	
2:00 298.0	152.1	0.0	0.0	351.6	348.6	348.5	342.9	3.2	5.7	6.1	6.5	395.9	373.1	
3:00 299.0	151.9	0.0	0.0	355.3	352.8	348.3	343.6	7.0	9.2	6.0	6.4	384.7	365.0	
4:00 298.0	152.0	0.0	0.0	356.2	353.7	349.3	345.3	6.9	8.4	6.1	6.5	383.8	366.1	
5:00 299.0	152.6	0.0	0.0	358.0	353.6	351.1	345.4	8.3	8.2	6.2	6.7	387.1	368.1	
6:00 299.0	152.4	0.0	0.0	362.1	355.6	353.9	347.2	8.3	8.4	6.3	6.8	384.8	368.2	
7:00 306.0	156.4	0.0	0.0	360.8	357.6	355.2	348.5	5.7	9.2	6.6	7.1	401.2	379.0	
8:00 322.0	165.3	0.0	0.0	362.4	357.5	355.0	349.1	7.4	3.4	6.9	7.6	424.9	396.9	
9:00 352.0	179.1	0.0	0.0	357.9	351.8	355.5	346.4	2.4	5.5	7.3	8.3	466.4	447.4	
10:00 350.0	179.7	0.0	0.0	348.0	346.7	346.9	340.7	1.1	6.0	7.3	7.5	476.0	431.5	
11:00 350.0	180.3	0.0	0.0	351.2	352.2	346.9	343.4	4.3	8.8	7.3	7.6	461.6	429.7	
12:00 351.0	180.8	0.0	0.0	360.1	358.9	353.2	348.0	6.9	10.9	7.5	7.8	465.1	427.9	
13:00 350.0	179.4	0.0	0.0	361.7	362.0	357.6	353.7	4.1	8.4	7.6	8.0	462.0	427.8	
14:00 350.0	180.2	0.0	0.0	362.0	362.4	356.9	354.1	5.1	8.3	7.5	7.9	462.0	428.4	
15:00 351.0	180.5	0.0	0.0	361.3	366.8	361.3	357.7	6.1	9.1	7.6	8.1	456.4	427.5	
16:00 350.0	180.5	0.0	0.0	373.0	371.3	365.6	362.1	7.3	9.2	7.7	8.2	454.0	423.6	
17:00 351.0	183.6	0.0	0.0	376.4	374.4	370.9	366.1	5.5	8.3	7.8	8.3	454.2	422.8	
18:00 355.0	185.0	0.0	0.0	362.0	358.8	361.5	356.2	0.5	2.5	7.9	8.2	466.7	436.4	
19:00 361.0	185.0	0.0	0.0	364.0	360.4	359.2	354.1	4.8	6.3	7.8	8.1	462.1	435.1	
20:00 361.0	185.5	0.0	0.0	367.0	362.2	360.2	355.1	6.8	7.1	8.1	8.5	477.9	440.2	
21:00 360.0	182.8	0.0	0.0	370.5	363.6	364.4	356.6	2.1	3.7	8.3	8.7	481.2	450.1	
22:00 361.0	183.4	0.0	0.0	364.8	358.2	362.7	354.6	1.0	3.7	8.0	8.3	473.0	442.1	
23:00 361.0	183.4	0.0	0.0	352.8	351.2	351.8	347.5	1.0	6.6	7.7	8.0	473.3	435.1	
24:00 361.0	184.1	0.0	0.0	355.6	353.8	350.7	347.2	5.0		7.8	8.0	473.5	433.3	

PHASE II

10/ 9/79 0131: 9

	1	2	3	4	5	6	7	8	9	10	11	12	13	14
1 INLET PART LOADING EAST														
2 INLET PART LOADING EAST														
3 INLET PART LOADING WEST														
4 INLET PART LOADING WEST														
5 OUTLET PART LOADING EAST														
6 OUTLET PART LOADING EAST														
7 OUTLET PART LOADING WEST														
8 OUTLET PART LOADING WEST														
9 PART REMOVAL EFF EAST														
10 PART REMOVAL EFF WEST														
11 INLET S02 EAST 1 HR AVG														
12 INLET S02 EAST 1 HR AVG														
13 INLET S02 WEST 1 HR AVG														
14 INLET S02 WEST 1 HR AVG														
15														
16														
17														
18														
	GR/SCF	LR/MBBTU	GR/SCF	LR/MBBTU	GR/SCF	LR/MBBTU	GR/SCF	LR/MBBTU	%	%	PPM	LR/MBBTU	PPM	LR/MBBTU
1100	0.6	0.00	0.2	0.00	-0.0073	0.0000	* 0.0000	0.0000	01.298	00.000	262.40	0.48	322.93	0.75
2100	0.2	0.00	-0.5	0.00	-0.0079	0.0000	* 0.0000	0.0000	04.419	00.000	255.76	0.46	323.51	0.75
3100	0.2	0.00	-0.5	0.00	-0.0091	0.0000	* 0.0000	0.0000	04.387	00.000	262.83	0.48	322.12	0.78
4100	0.2	0.00	-0.4	0.00	-0.0082	0.0000	* 0.0000	0.0000	04.216	00.000	260.74	0.47	323.16	0.75
5100	0.2	0.00	-0.5	0.00	-0.0047	0.0000	* 0.0000	0.0000	02.377	00.000	257.95	0.47	318.05	0.75
6100	0.3	0.00	0.9	0.00	-0.0105	0.0000	* 0.0000	0.0000	03.867	00.000	257.67	0.47	313.33	0.74
7100	1.2	0.00	1.1	0.00	-0.0110	0.0000	* 0.0000	0.0000	00.897	00.000	249.09	0.45	318.67	0.74
8100	2.6	0.00	6.1	0.01	-0.0237	0.0000	* 0.0000	0.0000	00.141	00.000	255.97	0.46	319.40	0.74
9100	2.7	0.00	5.9	0.01	-0.0227	0.0000	* 0.0000	0.0000	00.099	00.000	253.21	0.46	315.44	0.75
10100	0.5	0.00	0.6	0.00	-0.0052	0.0000	* 0.0000	0.0000	01.605	00.000	251.35	0.46	305.41	0.73
11100	0.4	0.00	-0.2	0.00	-0.0068	0.0000	* 0.0000	0.0000	01.764	00.000	250.10	0.45	305.37	0.72
12100	0.7	0.00	2.7	0.00	-0.0067	0.0000	* 0.0000	0.0000	00.970	00.000	251.37	0.46	320.45	0.75
13100	2.1	0.00	4.4	0.01	-0.0058	0.0000	* 0.0000	0.0000	00.275	00.000	257.13	0.47	331.98	0.77
14100	0.6	0.00	1.7	0.00	-0.0027	0.0000	* 0.0000	0.0000	00.431	00.000	256.22	0.47	325.38	0.76
15100	0.5	0.00	0.1	0.00	-0.0027	0.0000	* 0.0000	0.0000	00.566	00.000	253.03	0.46	320.33	0.75
16100	0.2	0.00	-0.3	0.00	-0.0027	0.0000	* 0.0000	0.0000	01.449	00.000	251.37	0.46	314.99	0.72
17100	3.6	0.00	8.9	0.01	-0.0027	0.0000	* 0.0000	0.0000	00.077	00.000	257.31	0.47	321.74	0.74
18100	1.3	0.00	2.2	0.00	-0.0027	0.0000	* 0.0000	0.0000	00.209	00.000	252.98	0.46	305.75	0.72
19100	0.7	0.00	1.1	0.00	-0.0027	0.0000	* 0.0000	0.0000	00.380	00.000	255.91	0.47	303.88	0.73
20100	0.3	0.00	-0.2	0.00	-0.0026	0.0000	* 0.0000	0.0000	00.874	00.000	250.24	0.45	289.03	0.72
21100	0.3	0.00	2.6	0.00	-0.0026	0.0000	* 0.0000	0.0000	00.345	00.000	242.02	0.44	287.70	0.70
22100	2.5	0.00	7.5	0.01	-0.0026	0.0000	* 0.0000	0.0000	00.106	00.000	261.80	0.48	292.70	0.73
23100	0.4	0.00	0.3	0.00	-0.0026	0.0000	* 0.0000	0.0000	00.606	00.000	261.89	0.48	287.23	0.70
24100	0.2	0.00	0.5	0.00	-0.0026	0.0000	* 0.0000	0.0000	01.184	00.000	257.99	0.47	287.51	0.68

B-6

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 3

10/ 9/79 0:32:16

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
1 OUTLET S02 EAST HR AVG																
2 OUTLET S02 EAST HR AVG																
3 OUTLET S02 WEST HR AVG																
4 OUTLET S02 WEST HR AVG																
5 S02 REM EFF EAST HR AVG																
6 S02 REM EFF WEST HR AVG																
7 INLET NOX EAST HR AVG																
8 INLET NOX EAST HR AVG																
9 INLET NOX WEST HR AVG																
10 INLET NOX WEST HR AVG																
11 OUTLET NOX EAST HR AVG																
12 OUTLET NOX EAST HR AVG																
13 OUTLET NOX WEST HR AVG																
14 OUTLET NOX WEST HR AVG																
15 NOX REMOVAL EFF EAST																
16 NOX REMOVAL EFF WEST																
17																
18																
100	159.6	0.39	* 61.7	0.15	18.17	80.54	276.3	0.36	376.9	0.63	177.3	0.31	* 157.0	0.27	13.69	57.57
200	164.6	0.43	* 67.7	0.16	7.87	78.53	230.5	0.30	312.6	0.52	153.3	0.29	* 129.7	0.22	4.84	57.62
300	178.2	0.44	* 52.5	0.13	7.51	83.35	227.0	0.30	298.3	0.50	154.0	0.27	* 108.9	0.19	7.91	62.71
400	185.3	0.46	* 49.7	0.12	2.51	84.30	224.3	0.29	296.6	0.50	150.9	0.27	* 99.6	0.17	7.73	65.69
500	183.7	0.46	* 130.5	0.31	1.82	58.14	226.1	0.29	303.4	0.51	152.0	0.27	* 173.4	0.30	7.36	41.69
600	194.1	0.48	* 152.8	0.37	-2.93	50.32	228.2	0.30	308.9	0.52	158.5	0.28	* 194.6	0.33	5.13	35.80
700	186.4	0.47	* 176.4	0.42	-3.97	43.42	232.0	0.30	313.5	0.52	159.0	0.29	* 214.6	0.37	4.79	30.01
800	195.7	0.48	* 155.6	0.37	-4.06	50.04	242.5	0.32	330.9	0.55	166.3	0.30	* 209.3	0.36	6.64	35.33
900	187.1	0.46	* 148.7	0.36	0.09	52.10	297.3	0.39	421.6	0.72	200.2	0.35	* 257.5	0.45	8.61	37.97
1000	183.4	0.45	* 137.9	0.33	1.32	54.09	293.5	0.38	422.0	0.72	198.3	0.35	* 257.5	0.45	8.61	37.97
1100	185.7	0.45	* 136.1	0.32	-0.12	54.69	280.9	0.37	408.8	0.69	190.7	0.34	* 247.4	0.42	8.48	38.25
1200	175.3	0.43	* 128.9	0.30	3.92	60.16	248.3	0.32	353.4	0.59	161.0	0.28	* 203.9	0.35	12.54	41.02
1300	163.9	0.41	* 110.7	0.25	11.91	65.85	246.6	0.32	349.6	0.58	155.7	0.28	* 191.8	0.31	12.73	46.79
1400	144.6	0.36	* 58.4	0.14	23.37	81.92	264.7	0.32	372.0	0.62	162.1	0.29	* 145.8	0.25	16.96	60.49
1500	132.0	0.33	* 11.2	0.03	28.82	96.46	248.0	0.32	355.9	0.59	144.7	0.26	* 84.9	0.14	20.39	75.47
1600	126.3	0.31	* 82.4	0.19	32.28	73.22	239.2	0.31	326.4	0.54	135.9	0.24	* 156.5	0.26	23.76	50.93
1700	124.9	0.31	* 110.0	0.26	34.30	64.98	240.7	0.31	336.3	0.55	134.1	0.24	* 187.3	0.32	24.59	42.94
1800	121.6	0.30	* 98.8	0.24	35.00	67.08	291.2	0.38	385.8	0.65	164.5	0.29	* 222.7	0.38	23.54	41.21
1900	124.4	0.30	* 103.2	0.25	36.17	65.94	302.6	0.39	396.9	0.67	173.3	0.30	* 232.6	0.40	24.49	40.24
2000	122.7	0.29	* 100.2	0.23	35.56	65.76	306.5	0.40	396.4	0.68	174.0	0.30	* 236.1	0.42	25.37	39.33
2100	118.7	0.28	* 122.5	0.30	35.35	56.45	311.0	0.41	402.4	0.70	177.2	0.30	* 264.5	0.47	24.92	32.77
2200	133.6	0.32	* 154.7	0.38	32.93	45.97	304.2	0.40	400.9	0.69	180.3	0.31	* 283.2	0.50	22.08	27.78
2300	142.5	0.34	* 167.3	0.40	28.38	42.69	342.1	0.45	446.9	0.76	212.4	0.36	* 328.8	0.57	18.29	25.09
2400	142.9	0.34	* 166.4	0.40	27.18	40.96	350.0	0.46	478.9	0.81	218.7	0.38	* 339.1	0.59	17.85	27.76

87

10/ 9/79 0:34:26

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 5

13 OUTLET DAMPER COMP#13 EAST
14 OUTLET DAMPER COMP#14 EAST

15 * YES IN SERVICE FOR THE HP
16 * NO OUT SERVICE FOR THE HP
17 *****
18 *****

1 OUTLET DAMPER COMP# 1 EAST
2 OUTLET DAMPER COMP# 2 EAST
3 OUTLET DAMPER COMP# 3 EAST
4 OUTLET DAMPER COMP# 4 EAST
5 OUTLET DAMPER COMP# 5 EAST
6 OUTLET DAMPER COMP# 6 EAST

7 OUTLET DAMPER COMP# 7 EAST
8 OUTLET DAMPER COMP# 8 EAST
9 OUTLET DAMPER COMP# 9 EAST
10 OUTLET DAMPER COMP#10 EAST
11 OUTLET DAMPER COMP#11 EAST
12 OUTLET DAMPER COMP#12 EAST

1

2

3

4

5

6

7

8

9

10

11

12

13

14

COMP	1	2	3	4	5	6	7	8	9	10	11	12	13	14
1:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
2:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
3:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
4:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
5:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
6:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
7:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
8:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
9:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
10:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
11:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
12:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
13:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
14:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
15:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
16:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
17:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
18:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
19:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
20:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
21:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
22:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
23:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
24:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES

8-9

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 4

10/ 9/79 0:33:21

1 INLET 02 EAST 1 HR AVG
2 INLET 02 WEST 1 HR AVG
3 OUTLET 02 EAST 1 HR AVG
4 OUTLET 02 WEST 1 HR AVG
5 PERCENT INCR 02 EAST 1 HR AVG
6 PERCENT INCR 02 WEST 1 HR AVG
7 STACK OPACITY 1HR AVG
8 STACK SO2 1HR AVG
9 STACK NOX 1HR AVG
10 STACK 02 1HR AVG
11 BYPASS DAMPER STATUS EAST
12 BYPASS DAMPER STATUS WEST
13 AMBIENT TEMP INST
14 BARO PRESS UNCORR INST
15 CALC ENERGY CONSUMPTION
16 AIR/CLOTH RATIO EAST 1HR AVG
17 AIR/CLOTH RATIO WEST 1HR AVG
18

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17
%	%	%	%	%	%	%	%	LR/ MMBTU	LR/ MMBTU	%	OPEN/ SHUT	OPEN/ SHUT	DEG F	IN HG	KWH	FT/ MIN	FT/ MIN
1:00 * 0.0	4.5	5.2	4.8	919.0	6.4	8.7	0.71	0.67	5.5	SHUT	SHUT	SHUT	70.1	26.21	2103.4	3.36	3.19
2:00 * 0.0	4.5	5.1	4.8	919.0	7.6	8.6	0.71	0.55	6.1	SHUT	SHUT	SHUT	68.9	26.20	1673.1	3.07	2.88
3:00 * 0.0	4.5	5.4	4.9	919.0	7.2	8.6	0.71	0.51	5.8	SHUT	SHUT	SHUT	69.4	26.21	1603.4	2.98	2.82
4:00 * 0.0	4.5	5.5	4.8	919.0	7.5	8.6	0.70	0.51	5.8	SHUT	SHUT	SHUT	62.4	26.20	1624.1	2.98	2.83
5:00 * 0.0	4.6	5.6	4.9	919.0	6.8	8.6	0.70	0.52	5.9	SHUT	SHUT	SHUT	62.0	26.21	1682.1	3.01	2.85
6:00 * 0.0	4.6	5.5	4.9	919.0	6.3	8.4	0.70	0.52	5.9	SHUT	SHUT	SHUT	61.0	26.21	1703.1	3.01	2.86
7:00 * 0.0	4.4	5.7	4.7	919.0	7.8	8.3	0.69	0.54	5.9	SHUT	SHUT	SHUT	63.3	26.21	1851.1	3.15	2.95
8:00 * 0.0	4.5	5.4	4.8	919.0	7.7	8.6	0.71	0.57	5.9	SHUT	SHUT	SHUT	61.8	26.21	2091.9	3.35	3.11
9:00 * 0.0	4.8	5.3	5.0	919.0	5.1	8.7	0.73	0.75	5.7	SHUT	SHUT	SHUT	61.9	26.21	2568.7	3.72	3.53
10:00 * 0.0	4.8	5.3	5.0	919.0	5.3	8.7	0.71	0.54	0.0	SHUT	SHUT	SHUT	66.9	26.21	2350.3	3.75	3.37
11:00 * 0.0	4.6	5.3	4.9	919.0	6.8	8.7	0.50	0.52	0.0	SHUT	SHUT	SHUT	74.3	26.21	2316.8	3.63	3.37
12:00 * 0.0	4.5	5.3	4.8	919.0	7.7	8.6	0.50	0.45	0.0	SHUT	SHUT	SHUT	87.4	26.21	2401.0	3.69	3.38
13:00 * 0.0	4.4	5.6	4.7	919.0	8.4	8.7	0.52	0.45	0.0	SHUT	SHUT	SHUT	91.3	26.21	2449.5	3.69	3.40
14:00 * 0.0	4.5	5.4	4.6	919.0	3.0	8.9	0.51	0.49	0.0	SHUT	SHUT	SHUT	93.5	26.22	2428.2	3.68	3.41
15:00 * 0.0	4.2	5.4	4.7	919.0	10.6	8.7	0.51	0.45	0.0	SHUT	SHUT	SHUT	96.0	26.21	2466.7	3.66	3.42
16:00 * 0.0	4.3	5.2	4.6	919.0	8.7	8.6	0.51	0.43	0.0	SHUT	SHUT	SHUT	96.9	26.21	2495.3	3.46	3.41
17:00 * 0.0	4.2	5.3	4.6	919.0	9.2	8.7	0.52	0.43	0.0	SHUT	SHUT	SHUT	96.7	26.21	2548.7	3.69	3.42
18:00 * 0.0	4.6	5.3	4.9	919.0	6.3	8.7	0.51	0.54	0.0	SHUT	SHUT	SHUT	94.1	26.21	2593.3	3.75	3.49
19:00 * 0.0	4.6	4.9	4.9	919.0	6.6	8.8	0.51	0.55	0.0	SHUT	SHUT	SHUT	87.8	26.21	2547.7	3.70	3.47
20:00 * 0.0	5.0	4.9	5.3	919.0	5.6	8.8	0.53	0.56	0.0	SHUT	SHUT	SHUT	70.9	26.21	2706.8	3.85	3.52
21:00 * 0.0	5.2	4.9	5.3	919.0	6.5	8.6	0.54	0.57	0.0	SHUT	SHUT	SHUT	74.2	26.21	2839.8	3.70	3.62
22:00 * 0.0	4.9	4.9	5.2	919.0	6.9	8.7	0.52	0.55	0.0	SHUT	SHUT	SHUT	71.4	26.21	2456.2	3.91	3.53
23:00 * 0.0	4.7	4.9	5.0	919.0	6.0	8.6	0.52	0.62	0.0	SHUT	SHUT	SHUT	69.4	26.20	2499.1	3.75	3.44
24:00 * 0.0	4.7	4.9	5.0	919.0	6.6	8.8	0.50	0.64	0.0	SHUT	SHUT	SHUT	69.5	26.22	2528.6	3.76	3.42

BZ

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 6

10/ 9/79 003520

13 OUTLET DAMPER COMP#27 WEST
14 OUTLET DAMPER COMP#28 WEST
15 *****
16 * YES IN SERVICE FOR THE HR*
17 * NO OUT SERVICE FOR THE HR*
18 *****

1 OUTLET DAMPER COMP#15 WEST
2 OUTLET DAMPER COMP#16 WEST
3 OUTLET DAMPER COMP#17 WEST
4 OUTLET DAMPER COMP#18 WEST
5 OUTLET DAMPER COMP#19 WEST
6 OUTLET DAMPER COMP#20 WEST

7 OUTLET DAMPER COMP#21 WEST
8 OUTLET DAMPER COMP#22 WEST
9 OUTLET DAMPER COMP#23 WEST
10 OUTLET DAMPER COMP#24 WEST
11 OUTLET DAMPER COMP#25 WEST
12 OUTLET DAMPER COMP#26 WEST

11

	1	2	3	4	5	6	7	8	9	10	11	12	13	14
COMP 15	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
1:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
2:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
3:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
4:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
5:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
6:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
7:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
8:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
9:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
10:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
11:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
12:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
13:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
14:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
15:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
16:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
17:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
18:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
19:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
20:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
21:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
22:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
23:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
24:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES

B-10

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER EAST INLET

3

ORSAT GAS ANALYSIS

	1	2	3
CO2	12.30	13.40	13.40
O2	9.00	8.30	8.06
N2	80.10	80.30	80.53
EXCESS AIR	50.08	41.84	39.91
DRY GAS MOL WT	30.25	30.30	30.39

PARTICULATE DATA
 BAROMETRIC PRESSURE
 NOZZLE AREA
 PILOT TUBE COEFFICIENT
 DUCT CROSS SECT MEA
 DUCT FLOW RATE
 SAMPLE WEIGHT
 SAMPLE VOLUME
 AVERAGE TEMPERATURE
 AVERAGE SOOT DELTA P
 AVERAGE DUCT FLOW RATE
 AVERAGE FUEL COAL
 AVERAGE HEAT RATE
 AVERAGE OPACITY

	1	2	3
IN HG	26.38	26.30	26.17
IN H2O	-17.53	-17.50	-16.20
FT2	360.2	379.5	385.2
IN H2O	0.950	0.950	0.850
FT2	246.0	210.0	246.0
MIN	123.40	112.82	121.27
FT3	231.40	231.40	231.40
GRAMS	9.0300	9.0300	9.0300
DEG R	54.13	54.13	54.13
IN H2O	0.82	0.84	0.83
MM	350.5	350.5	350.5
TON/HR	177.2	177.2	177.2
BTU/LB	8936	8936	8936
10**6 BTU/HR	3166.6	3166.6	3166.6
%	8.8	8.7	8.6

PARTICULATE RESULTS

SAMPLE VOL DRY STD COND
 PERCENT MULT ME
 WET STACK GAS MOL WT
 ABSOLUTE DUCT PRESS
 AVE DUCT VELOCITY
 DUCT FLOW DRY STD COND
 DUCT FLOW WET STD COND
 PART EMISS RATE
 PART EMISS RATE
 ISO KINETIC VARIATION

	1	2	3
FT3	99.573	99.573	106.087
VOL %	6.87	6.87	20.04
IN HG	28.00	28.00	28.00
FT3	62.38	62.38	62.38
IN HG	0.494	0.478	0.467
10**6 SCFM	0.921	0.934	0.928
10**6 SCFM	1.3993	1.3993	1.3993
GRAINS/SCF	4571.23	4571.23	4761.43
LB/HR	1.4436	1.4436	1.5235
LP/10**6 BTU	97.49	97.49	98.89

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER WEST INLET

6

DESAT GAS ANALYSIS

	A	S	6
CO2	13.20	13.43	13.35
O2	9.00	9.00	9.00
CO	80.90	80.63	80.65
POCF35 AIR	30.17	30.33	30.24
DRY GAS MOL WT	30.15	30.55	30.34

PARTICULATE DATA

BAROMETRIC PRESSURE	IN HG	26.30	26.17
STATIC PRESSURE	IN H2O	-17.30	-17.70
PISTON TUBE COEFFICIENT	FT2	0.3572E-03	0.3652E-03
DMCF	0.860	0.860	0.860
DUCT CROSS SECT AREA	FT2	0.974	0.974
TOTAL SAMPLING TIME	MIN	150	147
GAS SAMPLE VOL METER COND	FT3	122.83	121.07
MOISTURE COLLECTED	GRAMS	21.50	100.50
SAMPLE WEIGHT	GRAMS	81.50	146.0
Avg GAS METER TEMP	DEG R	572.9	505.3
Avg SO2 METER TEMP	DEG R	573.7	544.0
Avg SO2 METER DELTA P	IN H2O	0.83	0.80
Avg DELTA H	MM	2.00	2.38
Avg LOAD	TON/HR	350.5	350.5
Avg COAL FLOW RATE	BTU/LB	177.2	176.4
Avg BTU COAL	1000 BTU/HR	3174.7	3131.7
Avg HEAT RATE	1000 BTU/HR	3174.7	3131.7
Avg CAPACITY	1000 BTU/HR	3174.7	3131.7

PARTICULATE RESULTS

SAMPLE VOL DRY STD COND	FT3	97.508	103.394
PERCENT MOISTURE	VOL %	5.52	5.52
WET STACK GAS DRY STD	FT3	25.18	25.49
Avg DUCT VELOCITY	FT/SEC	62.83	61.69
DUCT FLOW DRY STD COND	1000 DSCFM	0.473	0.473
DUCT FLOW WET STACK COND	1000 DSCFM	0.927	0.927
PARTICULATE CONC	GRAINS/SCF	1.4130	1.4127
PART EMISS RATE	LB/HR	4527.62	4180.84
PART EMISS RATE	LB/HR	1.426	1.316
ISOKEINATIC VARIATION	LP/1000 BTU	94.71	101.24

SOUTHWESTERN PUBLIC SERVICE CO.
PARTICULATE TEST SUMMARY

TEST NUMBER	STACK	7	8	9
CRSAT GAS ANALYSIS				
CO2	X	18.72	19.57	13.20
CO	X	0.30	0.0	0.0
O2	X	6.30	6.03	6.26
N2	X	80.97	79.43	80.54
EXCESS AIR	X	41.78	40.41	41.78
DRY GAS MOL WT	X	30.29	30.57	30.36
PARTICULATE DATA				
BAROMETRIC PRESSURE	IN HG	26.24	26.15	26.03
STATIC PRESSURE	IN H2O	-0.89	-0.88	-0.80
NOZZLE AREA	FT2	.1444E-03	.1442E-03	.1473E-03
PILOT TUBE COEFFICIENT		0.870	0.870	0.870
DPVCF		3.973	3.973	3.973
DUCT CROSS SECT AREA	FT2	1.20	1.20	1.20
DUCT LENGTH	FT	54.03	54.03	54.75
GAS SAMPLE VOL MEAS COND		108.30	108.30	108.30
MOISTURE COLLECTED	GRAMS	0.0268	0.0308	0.0324
SAMPLE WEIGHT	DEGR	799.9	806.8	794.8
AVG STACK TEMP	DEGR	528.5	528.5	528.5
AVG GAS METER TEMP	IN H2O	1.2	1.2	1.2
AVG SO ROOT DELTA	IN H2O	1.2	1.2	1.2
AVG DELTA W	TON/HR	350.5	350.5	350.5
AVG COLD FLOW RATE	RTU/HR	177.2	176.4	176.8
AVG FTD COAL		8936	8555	8853
AVG HEAT RATE	10**6 BTU/HR	3166.6	3174.3	3131.3
AVG OPACITY	X	8.8	8.7	8.6
PARTICULATE RESULTS				
SAMPLE VOL DRY SED COND	FT3	42.271	44.709	45.694
PERCENT MOISTURE	VOL %	10.44	10.46	10.46
WET STACK GAS MJL AT		29.01	29.29	29.07
ABSOLUTE DUCT PRESS	IN HG	26.17	26.19	26.17
AVG DUCT VELOCITY	FT/SEC	8.55	8.55	8.55
DUCT FLOW DRY SED COND	10**6 BTU/HR	1.730	1.723	1.730
DUCT FLOW WET STACK COND	GRAMS/SEC	0.0098	0.0107	0.0109
PARTICULATE CONC	LB/10**6 BTU	62.44	65.54	68.29
PART EMISS RATE	LB/HR	0.0177	0.0207	0.0219
PART EMISS RATE	LB/10**6 BTU	51.40	54.93	57.07
ISOKINETIC VARIATION	X			

SOUTHWESTERN PUBLIC SERVICE CO.
PARTICULATE TEST SUMMARY

TEST NUMBER	WEST OUTLET	10	11	12
ORSAT GAS ANALYSIS				
CO ₂	X	12.73	12.74	12.63
CO	X	0.0	0.0	0.0
O ₂	X	6.13	6.30	6.36
N ₂	X	81.13	80.74	81.00
EXCESS AIR		40.14	43.64	42.34
DRY GAS MOL WT		30.22	30.30	30.28
PARTICULATE DATA				
BAROMETRIC PRESSURE	IN HG	26.38	26.33	26.17
STATIC PRESSURE	IN H ₂ O	-24.50	-25.80	-24.50
NOZZLE AREA	IN ²	0.860	0.860	0.860
PITOT TUBE COEFFICIENT	FT/SEC	31978	31978	31978
DMGT CROSS SECT AREA	FT ²	0.000	0.000	0.000
TOTAL SAMPLING TIME	MIN	14.5	14.5	14.5
GAS SAMPLE VOLUME	FT ³	67.1	62.79	98.61
MOISTURE COLLECTED	GRAMS	142.00	177.00	171.10
SAMPLE WEIGHT	GRAMS	0.0438	0.1111	0.1397
AVG STACK TEMP	DEG R	782.3	786.2	765.5
AVG GAS METER TEMP	DEG R	558.6	562.1	541.2
AVG SOOT METER DELTA P	IN H ₂ O	0.25	0.25	0.25
AVG DUCT AREA	IN ²	350.5	353.5	350.6
AVG COAL FLOW RATE	TON/HR	177.2	176.4	176.8
AVG FTD COAL	TON/HR	8936	8936	8936
AVG MEAT RATE	10000 BTU/HR	3166.6	3174.3	3131.3
AVG CAPACITY	%	8.8	8.7	8.6

B-14

PARTICULATE RESULTS				
SAMPLE VEL DRY STA COND	FT/SEC	58.120	70.337	78.038
PERCENT MOISTURE	VOL %	10.40	11.60	9.48
WET STACK GAS MOL WT		29.00	29.00	29.12
ABSOLUTE DUCT PRESS	IN HG	24.50	24.50	24.37
DUCT FLOW VELOCITY	FT/SEC	40.34	35.23	39.77
DUCT FLOW DRY STA COND	10000 BTU/HR	0.893	0.893	0.893
DUCT FLOW DRY STA COND	10000 BTU/HR	0.0116	0.0281	0.0276
PARTICULATE CONC	GRAMS/SEC	16.26	24.54	29.42
PART EMISS RATE	LB/10000 BTU	0.0114	0.0281	0.0286
PART EMISS RATE	LB/10000 BTU	105.13	104.34	116.85
ISO KINETIC VARIATION				

TEST NUMBER EAST OUTLET

PARTICULATE RESULTS

SAMPLE VOL	DRY ST	COND	FT3	62.018	5.540	6.170
PERCENT MULTIPLE			VOL %	25.52	10.51	10.54
NET STACK GAS MOL WT			IN HG	24.52	24.52	24.56
AVG DUCT PRESS			FT/SEC	39.54	37.47	38.18
AVG DUCT VELOCITY				0.44	0.40	0.42
DUCT FLUX DRY STD COND			1006.05CF4	0.375	0.329	0.345
DUCT FLUX WET STD COND			1006.05CF4	0.015	0.0119	0.0208
PARTICULATE CONC			GRAIN/CC	0.015	0.014	0.0208
PARTICULATE RATE			LB/HR	0.015	0.014	0.0208
HEAT EMISS RATE			B3/10006 BTU	0.12	0.11	0.12
HEAT EMISS RATE			%	56.43	56.43	56.43

B-15

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 2

3/24/80 10:23:25

	1	2	3	4	5	6	7	8	9	10	11	12	13	14
1 INLET PART LOADING EAST	GR/	LB/	GR/	LB/	GR/	LB/	GR/	LB/						
2 INLET PART LOADING EAST	SCF	MWRTU	SCF	MWRTU	SCF	MWRTU	SCF	MWRTU						
3 INLET PART LOADING WEST														
4 INLET PART LOADING WEST														
5 OUTLET PART LOADING EAST														
6 OUTLET PART LOADING EAST														
7 OUTLET PART LOADING WEST														
8 OUTLET PART LOADING WEST														
9 PART REMOVAL EFF EAST														
10 PART REMOVAL EFF WEST														
11 INLET S02 EAST 1 HR AVG														
12 INLET S02 EAST 1 HR AVG														
13 INLET S02 WEST 1 HR AVG														
14 INLET S02 WEST 1 HR AVG														
15														
16														
17														
18														
1100	GR/	LB/	GR/	LB/	GR/	LB/	GR/	LB/						
2100	SCF	MWRTU	SCF	MWRTU	SCF	MWRTU	SCF	MWRTU						
3100														
4100														
5100														
6100														
7100														
8100														
9100														
10100														
11100														
12100														
13100														
14100														
15100														
16100														
17100														
18100														

8-17

3/24/80 19:24:30

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 3

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
1 OUTLET S02 EAST 1 HR AVG																
2 OUTLET S02 EAST 1 HR AVG																
3 OUTLET S02 WEST 1 HR AVG																
4 OUTLET S02 WEST 1 HR AVG																
5 S02 REM EFF EAST 1 HR AVG																
6 S02 REM EFF WEST 1 HR AVG																
7 INLET NOX EAST 1 HR AVG																
8 INLET NOX EAST 1 HR AVG																
9 INLET NOX WEST 1 HR AVG																
10 INLET NOX WEST 1 HR AVG																
11 OUTLET NOX EAST 1 HR AVG																
12 OUTLET NOX EAST 1 HR AVG																
13 OUTLET NOX WEST 1 HR AVG																
14 OUTLET NOX WEST 1 HR AVG																
15 NOX REMOVAL EFF EAST																
16 NOX REMOVAL EFF WEST																
17																
18																
PPM	212.4	207.9	222.2	243.9	258.8	253.3	267.8	242.3	187.7	188.9	183.9	165.6	161.7	148.7	133.3	131.1
LB/MHTU	0.52	0.50	0.52	0.63	0.68	0.68	0.68	0.57	0.46	0.43	0.41	0.40	0.37	0.35	0.32	0.31
PPM	58.1	53.0	50.6	53.6	52.8	40.1	40.1	13.5	8.2	9.3	9.2	55.9	175.4	188.7	191.5	186.6
LB/MHTU	0.14	0.12	0.12	0.13	0.14	0.10	0.10	0.03	0.02	0.02	0.02	0.13	0.40	0.45	0.44	0.43
%	4.64	8.35	7.75	1.23	9.76	7.70	2.94	7.09	14.67	18.42	18.51	22.01	24.49	31.02	34.93	36.80
%	130.16	128.02	126.92	126.78	125.15	118.64	99.47	93.67	104.51	105.33	70.24	9.25	2.52	-0.41	-2.79	-2.30
PPM	244.6	265.8	290.3	191.0	133.8	142.5	149.6	229.7	272.3	256.7	274.4	279.4	290.8	287.1	299.0	297.2
LB/MHTU	0.44	0.47	0.49	0.36	0.26	0.28	0.32	0.40	0.49	0.42	0.49	0.48	0.49	0.50	0.52	0.51
PPM	278.8	312.0	336.1	203.2	121.3	136.6	162.0	253.8	330.0	297.3	341.4	340.1	352.4	352.1	386.9	388.1
LB/MHTU	0.46	0.52	0.58	0.36	0.22	0.25	0.29	0.44	0.56	0.51	0.58	0.58	0.59	0.59	0.65	0.65
PPM	245.4	259.0	284.0	211.6	167.7	174.9	199.7	241.4	254.4	233.8	238.6	243.1	250.4	241.0	249.5	246.6
LB/MHTU	0.43	0.45	0.47	0.39	0.32	0.34	0.36	0.44	0.44	0.38	0.41	0.41	0.42	0.41	0.43	0.42
PPM	2.8	12.2	16.8	10.3	4.0	9.8	45.3	74.9	9.5	20.5	91.1	274.6	289.3	316.4	352.1	349.2
LB/MHTU	0.00	0.02	0.03	0.02	0.01	0.02	0.08	0.13	0.02	0.04	0.16	0.48	0.50	0.54	0.60	0.59
%	1.31	3.91	9.13	94.88	96.66	92.72	71.68	69.65	102.91	107.03	13.21	15.31	15.80	18.07	18.35	18.14
%	99.00	96.05	94.93	94.88	96.66	92.72	71.68	69.65	102.91	107.03	13.21	15.31	15.80	18.07	18.35	18.14
%	1.31	3.91	9.13	94.88	96.66	92.72	71.68	69.65	102.91	107.03	13.21	15.31	15.80	18.07	18.35	18.14
%	99.00	96.05	94.93	94.88	96.66	92.72	71.68	69.65	102.91	107.03	13.21	15.31	15.80	18.07	18.35	18.14

B-18

3/24/80 19:25:35

B-19

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 5

3/24/80 19:26:39

1-OUTLET-DAMPER-COMP# 1-EAST																	
2-OUTLET-DAMPER-COMP# 2-EAST																	
3-OUTLET-DAMPER-COMP# 3-EAST																	
4-OUTLET-DAMPER-COMP# 4-EAST																	
5-OUTLET-DAMPER-COMP# 5-EAST																	
6-OUTLET-DAMPER-COMP# 6-EAST																	
7-OUTLET-DAMPER-COMP# 7-EAST																	
8-OUTLET-DAMPER-COMP# 8-EAST																	
9-OUTLET-DAMPER-COMP# 9-EAST																	
10-OUTLET-DAMPER-COMP# 10-EAST																	
11-OUTLET-DAMPER-COMP# 11-EAST																	
12-OUTLET-DAMPER-COMP# 12-EAST																	
13-OUTLET-DAMPER-COMP# 13-EAST																	
14-OUTLET-DAMPER-COMP# 14-EAST																	
15-*****																	
16- * YES IN SERVICE FOR THE HR*																	
17- * NO OUT SERVICE FOR THE HR*																	
18-*****																	
1	2	3	4	5	6	7	8	9	10	11	12	13	14				
COMP	COMP	COMP	COMP	COMP	COMP	COMP	COMP	COMP	COMP	COMP	COMP	COMP	COMP				
1:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
2:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
3:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
4:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
5:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
6:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
7:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
8:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
9:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
10:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
11:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
12:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
13:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
14:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
15:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
16:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
17:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
18:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES

FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 6

B-21

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER	EU1 1	EU2 2	EU3 3
URSAT GAS ANALYSIS			
CO2	X	14.00	13.93
CO	X	0.0	0.0
O2	X	0.0	0.0
N2	X	80.93	81.06
EXCESS AIR	X	30.86	30.86
DRY GAS MOL WT	X	30.44	30.42
PARTICULATE DATA			
BAROMETRIC PRESSURE	IN HG	26.07	26.12
STATIC PRESSURE	IN H2O	26.50	26.50
NOZZLE AREA	FT2	0.0012	0.0012
PITOT TUBE COEFFICIENT		0.840	0.840
DMCF		0.980	0.980
DUCT CROSS SECT AREA	FT2	369.0	369.0
DUCT LENGTH	FT	147	147
GAS SAMPLE VOLUME	MIN	168.72	168.72
MOISTURE COLLECTED	GRAMS	268.00	268.00
SAMPLE WEIGHT	GRAMS	271.2	271.2
AVG STACK TEMP	DEG R	548.1	548.1
AVG GAS METER TEMP	IN H2O	0.53	0.53
AVG DUCT METER TEMP	IN H2O	0.53	0.53
AVG DUCT AREA	FT2	369.0	369.0
AVG DUCT FLOW RATE	TON/HR	350.8	350.8
AVG BTU COAL	BTU/LB	187.7	187.7
AVG HEAT RATE	10**6 BTU/HR	331.2	331.2
AVG OPACITY	%	2.3	2.3

PARTICULATE RESULTS

SAMPLE VOL DRY STD COND	FT3	140.106	58.190	58.266
PERCENT MOISTURE		21.14	21.14	21.14
WET STACK GAS MOL WT		23.55	23.55	23.55
ABSOLUTE DUCT PRESS	IN HG	23.97	23.97	23.97
DUCT FLOW VELOCITY	FT/SEC	39.47	39.47	39.47
DUCT FLOW WET STACK COND	10**6 DSCFM	0.445	0.445	0.445
DUCT FLOW WET STACK COND	GRAMS/SEC	0.074	0.074	0.074
PARTICULATE CONC	LB/10**6 HTU	0.0081	0.0131	0.0113
PART EMISS RATE		2.072	3.865	3.350
ISOKINETIC VARIATION	%	98.77	95.16	92.69

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER	ED4 4	MOL 5	MOL 6
OBSAT GAS ANALYSIS			
CO2	13.73	12.87	11.53
CO	9.30	8.16	5.56
O2	80.97	80.96	80.90
N2	32.97	40.53	35.24
EXCESS AIR	30.41	30.31	30.39
DRY GAS MOL WT			
PARTICULATE DATA			
BAROMETRIC PRESSURE	IN HG	26.07	26.18
STATIC PRESSURE	IN H2O	-27.20	-27.20
NOZZLE AREA	FT2	.81694E-03	.40952E-03
PITOT TUBE COEFFICIENT		0.860	0.860
DUCT CROSS SECT AREA	FT2	0.991	0.991
TOTAL SAMPLING TIME	MIN	369.0	369.0
MOISTURE COLLECTED	GRAMS	150.85	71.46
GAS SAMPLE VOL METER COND	GRAMS	249.00	153.00
SAMPLE WEIGHT	GRAMS	0.1005	0.1171
Avg Gas Meter Temp	DEG R	775.2	775.2
Avg SO H2O DELTA P	IN H2O	546.0	550.0
Avg Delta H	MM	0.49	0.23
Avg Coal Flow Rate	TON/HR	350.8	345.7
Avg BTU Coal	MM	186.5	187.7
Avg Heat Rate	MM	8932	8872
Avg Opacity	10**6 BTU/HR	3331.0	3331.4
		2.3	1.9
PARTICULATE RESULTS			
SAMPLE VOL DRY STD COND	FT3	56.494	59.508
PERCENT MOISTURE	WT	12.17	10.83
RESOLUTE DUCT PRESS	IN HG	28.90	29.27
Avg Duct Velocity	FT/SEC	23.72	24.07
Duct Flow Dry STD COND	10**6 DSCFM	41.56	38.22
Duct Flow Wet Stack Cond	10**6 CFM	0.425	0.401
Particulate Conc	GRAINS/SCF	0.0020	0.0020
Part Emiss Rate	LB/10**6 BTU	0.2118	0.198
Isokinetic Variation		0.0065	0.0126
		93.48	91.07

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER		MO3 7	MO4 8
ORSAT GAS ANALYSIS			
CO2	X	13.40	13.56
CO	X	0.0	0.0
O2	X	5.37	5.53
N2	X	81.23	80.90
EXCESS AIR	X	33.41	34.98
DRY GAS MOL WT	X	30.36	30.39
PARTICULATE DATA			
BAROMETRIC PRESSURE	IN HG	26.12	25.91
STATIC PRESSURE	IN H2O	-27.20	-27.20
NOZZLE AREA	IN ²	.80952E-03	.80653E-03
PITOT TUBE COEFFICIENT		0.860	0.860
UGMCF CROSS SECT AREA	FT2	0.991	0.991
DUCT SAMPLE VOLUME	MIN	369.0	369.0
DUCT SAMPLE VOLUME	FT3	1.97	1.97
MOISTURE COLLECTED	GRAMS	131.50	197.50
SAMPLE WEIGHT	DEG R	0.0756	0.0905
AVG STACK TEMP	DEG R	793.0	782.6
AVG GAS METER TEMP	DEG R	568.3	535.2
AVG SO ROOT DELTA P	IN H2O	0.48	0.50
AVG DELTA P	IN H2O	0.475	0.475
AVG FLOW RATE	TON/HR	350.2	345.0
AVG BTU COAL	BTU/LB	8866.2	8840.0
AVG HEAT RATE	10**6 BTU/HR	3313.6	3264.9
AVG DENSITY	X	2.1	2.2
PARTICULATE RESULTS			
SAMPLE VOL DRY STD COND	FT3	53.922	68.097
PERCENT MOISTURE	VOL X	10.11	12.05
NET STACK GAS MOL WT		29.08	28.90
ABSOLUTE DUCT PRESS	IN HG	24.12	23.91
AVG DUCT VELOCITY	FT/SEC	37.85	39.05
DUCT FLOW DRY STD COND	10**6 DSCFM	0.403	0.460
DUCT FLOW WET STD COND	10**6 CFM	0.839	0.860
DUCT FLOW WET STACK COND	GRAINS/SCF	0.826	0.8205
PARTICULATE CONC	LB/HR	0.962	65.04
PART EMISS RATE	LB/10**6 BTU	0.0207	0.0199
ISOKINETIC VARIATION	X	81.97	101.89

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PHASE IV

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106 INLET S02 WEST 1 HR AVG

107 INLET S02 WEST 1 HR AVG

PHASE IV

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FAN-10 FILTER SYSTEM SUMMARY LOG - PAGE 3

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
1 INLET NOX EAST	13.00	231.3	2.55	136.6	0.43	998.23	2.70	0.01	171.9	0.30	234.2	0.40	199.9	0.35	235.50	-17.69
2 INLET NOX WEST	13.00	225.7	0.92	147.7	0.47	939.21	0.55	0.00	167.3	0.30	230.3	0.33	195.3	0.35	645.20	-19.34
3 INLET NOX EAST	13.00	203.9	0.47	185.5	0.45	653.09	-1.25	0.01	180.3	0.33	234.7	0.39	211.5	0.38	492.84	-14.22
4 INLET NOX WEST	13.00	202.6	0.47	185.4	0.47	555.68	-6.74	0.01	191.3	0.34	239.9	0.40	213.1	0.38	405.34	-13.26
5 INLET NOX EAST	13.00	223.0	0.52	183.8	0.51	690.64	-9.47	0.00	173.6	0.31	244.0	0.41	215.5	0.40	555.60	-27.40
6 INLET NOX WEST	13.00	230.4	0.59	251.8	0.53	936.49	-14.93	0.00	166.2	0.12	206.3	0.33	187.3	0.34	622.01	185.98
7 INLET NOX EAST	13.00	226.9	0.52	206.1	0.51	801.71	-12.37	0.00	196.6	0.35	256.5	0.42	230.0	0.42	510.70	-68.32
8 INLET NOX WEST	13.00	232.0	0.53	181.4	0.45	888.63	-2.91	0.00	211.2	0.37	258.0	0.42	224.8	0.41	779.91	-8.55
9 INLET NOX EAST	13.00	227.2	0.52	178.7	0.44	901.14	4.47	0.00	215.5	0.38	248.7	0.40	222.1	0.40	724.11	-5.04
10 INLET NOX WEST	13.00	214.5	0.49	162.3	0.42	616.44	6.34	0.00	228.6	0.40	243.8	0.40	219.8	0.39	751.91	2.23
11 INLET NOX EAST	13.00	199.1	0.45	162.0	0.42	557.35	11.67	0.00	231.9	0.41	239.6	0.39	224.0	0.40	777.49	1.95
12 INLET NOX WEST	13.00	199.3	0.41	165.2	0.41	259.80	12.37	0.00	237.6	0.42	234.7	0.38	225.5	0.40	756.73	3.61
13 INLET NOX EAST	13.00	152.3	0.37	163.2	0.40	116.89	12.65	0.01	243.3	0.42	229.5	0.38	231.1	0.41	856.46	3.39
14 INLET NOX WEST	13.00	151.8	0.34	163.1	0.40	790.23	17.03	0.00	240.8	0.44	227.5	0.37	235.0	0.42	840.75	4.16
15 INLET NOX EAST	13.00	135.6	0.31	166.2	0.41	710.67	18.70	0.00	261.0	0.44	233.7	0.36	244.1	0.43	472.55	1.04
16 INLET NOX WEST	13.00	151.0	0.35	167.1	0.41	010.96	13.94	0.00	241.3	0.42	225.2	0.36	232.5	0.41	352.04	2.10
17 INLET NOX EAST	13.00	252.8	0.58	176.0	0.43	997.50	3.54	0.00	229.3	0.39	251.9	0.43	211.9	0.37	522.24	6.08
18 INLET NOX WEST	13.00	302.7	0.69	172.2	0.43	454.98	-4.03	0.00	240.8	0.43	283.5	0.46	214.5	0.38	070.03	12.45
19 INLET NOX EAST	13.00	294.1	0.69	175.4	0.42	639.62	-7.23	0.00	266.6	0.45	287.1	0.49	225.7	0.39	773.18	14.21
20 INLET NOX WEST	13.00	297.9	0.69	171.2	0.42	665.68	-14.91	0.00	259.4	0.45	291.1	0.48	228.0	0.40	713.90	10.78
21 INLET NOX EAST	13.00	264.7	0.66	182.3	0.45	257.20	-23.05	0.01	231.5	0.40	274.8	0.45	223.7	0.38	621.15	1.37
22 INLET NOX WEST	13.00	327.5	0.81	205.6	0.50	233.35	-21.63	0.00	161.0	0.28	255.5	0.45	184.4	0.32	527.81	-14.46

B-26

B-27

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FABRIC FILTER SYSTEM SUMMARY LOG - PAGE 6

	1	2	3	4	5	6	7	8	9	10	11	12	13	14
1	OUTLET DAMPER COMP#15 WEST													
2	OUTLET DAMPER COMP#16 WEST													
3	OUTLET DAMPER COMP#17 WEST													
4	OUTLET DAMPER COMP#18 WEST													
5	OUTLET DAMPER COMP#19 WEST													
6	OUTLET DAMPER COMP#20 WEST													
7	OUTLET DAMPER COMP#21 WEST													
8	OUTLET DAMPER COMP#22 WEST													
9	OUTLET DAMPER COMP#23 WEST													
10	OUTLET DAMPER COMP#24 WEST													
11	OUTLET DAMPER COMP#25 WEST													
12	OUTLET DAMPER COMP#26 WEST													
13	OUTLET DAMPER COMP#27 WEST													
14	OUTLET DAMPER COMP#28 WEST													
15	*****													
16	YES IN SERVICE FOR THE 12*													
17	*NO OUT SERVICE FOR THE 12*													
18	*****													
19:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
20:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
21:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
22:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
23:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
24:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
25:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
26:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
27:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
28:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
29:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
30:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
31:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
32:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
33:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
34:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
35:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
36:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
37:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
38:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
39:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
40:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
41:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
42:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
43:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
44:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
45:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
46:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
47:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
48:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
49:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
50:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
51:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
52:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
53:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
54:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
55:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
56:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
57:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
58:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
59:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES
60:00	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES

B-29

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER

E1 E12 E13

ORSAT GAS ANALYSIS

	E1	E12	E13
CO ₂	14.10	13.96	14.73
CO	0.00	0.00	4.40
O ₂	81.00	81.16	80.46
N ₂	20.90	29.37	29.19
EXCESS AIR	30.45	30.43	30.55
DRY GAS MOL WT			

PARTICULATE DATA

	E1	E12	E13
BAROMETRIC PRESSURE	26.44	26.41	26.42
STATIC PRESSURE	-17.70	-17.70	-17.70
NOZZLE AREA	0.637	0.637	0.637
PITOT TUBE COEFFICIENT	1.001	1.001	1.001
DOWN DRAUGHT	246.0	246.0	246.0
CROSS SECT AREA	187.	187.	187.
TOTAL SAMPLING TIME	110.02	110.82	117.7
MOISTURE COLLECTED	297.00	180.00	291.00
GAS SAMPLE VCL METER COND	92.14	111.10	81.110
SAMPLE WEIGHT	554.8	557.9	561.1
AVG GAS METER TEMP	55.79	55.05	56.11
AVG SOL ROOM DELTA P	0.79	0.78	0.79
AVG LEAK	1.69	1.66	1.69
AVG COAL FLOW RATE	385.3	389.5	389.5
AVG FTU COAL	184.3	195.7	190.8
AVG FEAT RATE	801.1	3259.1	3281.1
AVG DPACTV	8.3	8.3	8.1

PARTICULATE RESULTS

	E1	E12	E13
SAMPLE VCL DRY STD COND	93.013	92.994	92.754
PERCENT MOISTURE	3.62	3.62	3.75
WET STACK GAS MCL WT	25.18	25.94	28.95
ABSOLUTE DUCT PRESS	59.94	59.66	60.29
AVG DUCT VELOCITY	0.415	0.411	0.414
DUCT FLOW WET STACK COND	1046 DSCFM	1046 DSCFM	1046 DSCFM
PARTICULATE CONC	1.6447	1.6875	1.6875
PART EMISS RATE	4684.30	5360.62	3863.17
PART EMISS RATE	1.453	1.453	1.453
ISO KINETIC VARIATION	97.52	98.83	97.97

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER		M1	M2	M3
		4	5	6
UNSAT GAS ANALYSIS				
CO ₂	X	14.26	13.25	13.40
O ₂	X	6.69	9.92	9.90
N ₂	X	80.67	80.82	80.70
EXCESS AIR	X	39.68	38.44	38.30
DRY GAS MOL WT	X	30.37	30.36	30.38
PARTICULATE DATA				
BAROMETRIC PRESSURE	IN HG	26.44	26.41	26.42
STATIC PRESSURE	IN H ₂ O	-19.00	-19.00	-19.40
NOZZLE AREA	FT ²	.3830E-03	.3830E-03	.3830E-03
PITOT TUBE COEFFICIENT		0.845	0.845	0.845
DIFF. CROSS SECT. AREA	FT ²	0.997	0.997	0.997
DIFF. PRESS. TIME	FT	24.0	24.0	24.0
TOTAL SAMPLE VOL	FT ³	11.7	11.8	11.7
GAS SAMPLE VOL METER COND	FT ³	11.7	11.8	11.7
MOISTURE COLLECTED	GRAMS	243.00	243.00	243.00
SAMPLE WEIGHT	GRAMS	12.0332	10.2895	12.0332
AVG GAS METER TEMP	DEG F	788.2	793.4	788.2
AVG GAS METER TEMP	DEG C	554.7	559.1	559.9
AVG ROOT DELTA P	IN H ₂ O	0.80	0.81	0.80
AVG LEAKAGE	IN H ₂ O	2.50	2.50	2.50
AVG COAL FLOW RATE	TON/HR	18.3	18.3	18.3
AVG ETO COAL	BTU/LB	8113	8113	8113
AVG FERT RATE	1000 BTU/HR	3247.7	3250.4	3247.7
AVG CAPACITY	X	8.3	8.3	8.1
PARTICULATE RESULTS				
SAMPLE VOL DRY STD COND	FT ³	96.407	95.267	96.181
PERCENT MOISTURE	W/L	12.17	12.27	12.27
NET STACK GAS MOL WT		28.66	28.64	28.64
ABSOLUTE DUCT PRESS	IN HG	25.04	25.01	25.07
AVG DUCT VELOCITY	FT/SEC	60.18	61.00	60.82
AVG DUCT FLOW	1000 SCFM	0.437	0.439	0.441
DUCT FLOW DRY STD COND	1000 SCFM	1.088	0.900	0.898
PARTICULATE CONC	GRAINS/SCF	1.088	1.088	1.088
PART EMISS RATE	1000 BTU/HR	630.22	543.69	478.37
PART EMISS RATE	1000 BTU/HR	1.5567	1.5691	1.5567
ISOKEINEL VARIATION	X	98.40	98.11	98.38

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER	ST1	ST2	ST3
	7	8	9
ORSAT GAS ANALYSIS			
CO2	13.26	13.36	13.38
O2	0.00	0.00	0.00
N2	85.61	85.64	85.61
EXCESS AIR	30.06	30.06	30.10
CRY GAS MUL WT	30.35	30.36	30.39
PARTICULATE DATA			
BAROMETRIC PRESSURE	26.34	26.31	26.32
NOZZLE AREA	0.50	0.50	0.50
PITOT TUBE COEFFICIENT	0.971	0.971	0.971
DUCT CROSS SECT AREA	120.0	120.0	120.0
TOTAL SAMPLE VOLUME	53.49	56.89	57.81
MOISTURE COLLECTED	130.90	139.30	138.59
SAMPLE WEIGHT	0.0465	0.0503	0.0489
AVG GAS METER TEMP	51.5	54.1	54.1
AVG SQ ROOT DELTA P	1.06	1.07	1.09
AVG DELTA H	1.10	1.14	1.15
AVG CUL FLOW RATE	34.5	34.5	34.5
AVG FLOW RATE	184.3	185.7	185.8
AVG FLOW RATE	681.1	671.1	671.1
AVG FLAT RATE	3247.7	3258.5	3261.1
AVG CAPACITY	6.3	6.3	6.1
PARTICULATE RESULTS			
SAMPLE VOL DRY STD COND	45.03	47.951	48.369
PERCENT MOISTURE	12.07	12.20	12.20
WET STACK GAS MUL A1	28.66	28.66	28.66
ABSOLUTE DUCT PRESS	0.002	0.003	0.003
AVG DUCT VELOCITY	0.893	0.893	0.893
DUCT FLOW DRY STD COND	1.653	1.653	1.653
DUCT FLOW WET COND	0.0161	0.0094	0.0150
GRAINS/SCF	131.11	58.50	96.95
PART EMISS RATE	0.0311	0.0190	0.0295
PART EMISS RATE	89.60	95.71	95.66
ISOKINETIC VARIATION			

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER

E11 E12 E13

URSAT GAS ANALYSIS

CO2	X	12.16	14.40	17.02
O2	X	6.98	4.84	0.30
N2	X	80.66	80.76	80.68
EXCESS AIR		48.82	29.33	42.00
DRY GAS MOL WT		30.26	30.50	30.34

PARTICULATE DATA

BAROMETRIC PRESSURE	IN HG	29.29	26.21	26.17
STATIC PRESSURE	IN H2O	-17.70	-16.30	-16.30
NOZZLE AREA		.38300E-03	.38300E-03	.38300E-03
PITOT TUBE COEFFICIENT	FT2	0.802	0.802	0.802
DMCF	FT2	0.987	2.987	2.987
DUCT CROSS SECT AREA	FT2	21.70	11.70	11.70
DUCT FLOW TIME	FT	126.53	115.24	115.11
WET SAMPLE METER COND	GRAMS	67.50	287.00	248.00
MOISTURE COLLECTED	GRAMS	0.921	11.645	11.645
SAMPLE WEIGHT	DEG R	817.8	817.8	817.8
AVG STACK TEMP	IN H2O	570.8	565.6	522.4
AVG S3 ROOT DELTA P	IN H2O	9.61	1.87	1.87
AVG DELTA H	IN H2O	355.2	355.0	356.0
AVG COAL FLOW RATE	TON/HR	189.1	187.9	187.6
AVG BTU COAL	BTU/HR	8736	8774	8777
AVG HEAT RATE	1000 BTU/HR	3304.8	3298.1	3293.6
AVG CAPACITY		2.1	2.4	2.5

PARTICULATE RESULTS

SAMPLE VOL DRY STD COND	FT3	102.035	93.463	92.874
PERCENT MOISTURE	VOL %	3.03	12.67	11.20
NET STACK GAS MOL WT		29.09	28.01	28.09
ABSOLUTE DUCT PRESS	IN HG	24.98	25.01	24.97
AVG DUCT VELOCITY	FT/SEC	57.83	0.46	0.412
DUCT FLOW DRY STD COND	1000 SCFH	0.894	0.481	0.487
DUCT FLOW WET COND	1000 SCFH	814.6	1.928	1.3810
WET FLOW CORP	GRAMS/SEC	509.22	5679.04	4581.36
PART EMISS RATE	LH/HR	0.1541	1.7219	1.3910
ISO KINETIC VARIATION	LH/1000 BTU	99.59	100.34	98.42

B-32

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER	W1	W2	W3
	4	3	0
ORSAT GAS ANALYSIS			
CO2	X	14.00	13.26
CO	X	0.0	0.0
O2	X	5.20	6.24
N2	X	80.80	80.59
EXCESS AIR	X	32.24	41.02
DRY GAS MOL WT		30.45	30.37
PARTICULATE DATA			
BAROMETRIC PRESSURE	IN HG	26.28	26.21
STATIC PRESSURE	IN H2O	-19.03	-17.00
NOZZLE AREA	FT2	3.7730E-03	3.7730E-03
PITOT TUBE COEFFICIENT		0.824	0.824
DUCT CROSS SECT AREA	FT2	340.6	240.6
DUCT FLOW VELOCITY	FT/SEC	192	147
TOTAL SAMPLING TIME	MIN	118.54	112.11
GAS SAMPLE VOL METER COND	GRAMS	306.00	273.50
MOISTURE COLLECTED	GRAMS	14.2732	11.0003
SAMPLE WEIGHT	GRAMS	821.1	813.1
AVG STACK TEMP	DEG R	577.3	577.0
AVG GAS METER TEMP	DEG R	577.3	577.0
AVG DUCT TEMP	DEG R	248	145
AVG DUCT WIND DELTA P	IN H2O	255.2	350.0
AVG LEAD FLOW RATE	TON/HR	199.1	187.0
AVG COAL FLOW RATE	TON/HR	873.6	877.4
AVG BTU COAL	BTU/LB	3304.2	3293.1
AVG HEAT RATE	10**6 BTU/HR	241	244
AVG OPACITY	X		
PARTICULATE RESULTS			
SAMPLE VOL DRY STD COND	FT3	95.949	91.142
PERCENT MOISTURE		13.04	12.41
NET STACK GAS MOL WT		28.82	28.94
ABSOLUTE DUCT PRESS	IN HG	24.98	24.97
AVG DUCT VELOCITY	FT/SEC	0.417	0.413
DUCT FLOW VELOCITY	10**6 DSCM	0.926	0.870
DUCT FLOW WET STACK COND	GRAMS/SCF	2.2993	1.5081
PARTICULATE CONC	LB/10**6 BTU	7072.36	4310.18
PART EMISS RATE		2.1400	1.8522
PART EMISS VARIATION		98.86	97.97
ISOKINETIC VARIATION			97.35

SOUTHWESTERN PUBLIC SERVICE CO.

PARTICULATE TEST SUMMARY

TEST NUMBER

S1 7 S2 8 S3 9

URSAY GAS ANALYSIS

CO2	X	12.50	13.58	13.35
CO	X	0.0	0.0	0.0
O2	X	6.69	5.85	5.97
N2	X	80.80	80.56	80.69
EXCESS AIR	X	45.69	37.94	38.59
DRY GAS MOL WT		30.27	30.41	30.37

PARTICULATE DATA

BAROMETRIC PRESSURE	IN HG	26.18	26.11	26.07
STATIC PRESSURE	IN H2O	-1.00	-0.95	-0.97
PITOT TUBE COEFFICIENT	IN FT2	.14190E-03	.14190E-03	.14190E-03
DMCF	FT2	0.780	0.780	0.780
TOTAL CROSS SECT AREA	FT2	0.949	0.949	0.949
GAS SAMPLE VOL METER COND	FT3	31.25	31.25	31.25
MOISTURE COLLECTED	GRAMS	57.01	46.60	99.36
SAMPLE STACK TEMP	DEG R	132.50	114.00	225.90
AVG GAS METER TEMP	DEG R	825.53	816.71	927.8
AVG SO HORT DELTA P	IN H2O	1.19	1.00	1.08
AVG DELTA H	IN H2O	1.19	1.00	1.01
AVG LOAD	TON/HR	355.2	355.0	356.0
AVG COAL FLOW RATE	BTU/LB	189.1	187.9	187.6
AVG BTU CHATE	BTU/HR	8736	8774	8777
AVG CAPACITY	10**6 BTU/HR	3304.8	3298.1	3293.6
		2.1	2.4	2.5

PARTICULATE RESULTS

SAMPLE VOL DRY STD COND	FT3	45.870	37.860	79.597
PERCENT DRY STD COND	VL X	12.01	12.45	11.82
PERCENT DRY STD COND	VL X	28.80	28.86	28.00
ABSOLUTE DUCT PRESS	FT/SEC	26.11	26.09	25.01
DUCT FLOW DRY STD COND	10**6 DSCFM	82.59	0.775	0.753
DUCT FLOW WET STACK COND	10**6 DSCFM	1.868	1.573	1.529
PARTICULATE CONC	LB/10**6 BTU	0.0152	0.0192	0.0082
PART EMISS RATE	LB/10**6 BTU	104.62	121.91	53.44
PART EMISS RATE	LB/10**6 BTU	0.0317	0.0370	0.0162
ISO KINETIC VARIATION		11.16	97.84	99.41

Appendix C

Field Notes

Representative Field Notes
Phase III

Test Report for March, 1980, EPA Baghouse Study

3/24/80

1. West outlets SO₂ impingers being transferred forward, bad check valve
2. Electrical problem with hot box-east outlet. Rheostat in control box replaced.
3. Leak check problem on east outlet initially.
4. Hot box/cold box connector only are caused glassware to be washed each time (stack).
5. Hot box on east inlet hit conduit for pilot baghouse during SO₂ sample #6, causing it to be damaged and caused the probe liner to partially come out of probe. After hot box was replaced, an acceptable leak check could not be obtained. After delaying other stations for about an hour. It was decided to proceed without the east inlet.
6. While the east inlet was having problems, the west outlet could not obtain an acceptable leak rate. At the same time the east inlet was not being sampled, it was decided not to sample the west outlet. While other stations were sampling it was discovered that a faulty connection on the umbilical cord was the problem. This was tightened and the problem was solved.
7. At about 6:30, it was noticed on the opacity chart that an intermittent spike was developing. This aroused suspicion of a broken bag.

3/25/80

1. The unit was confirmed to have a broken bag in compartment #23. After inspection two broken bags were discovered. These bags were removed and the thimbles were blanked off.
2. The unit momentarily tripped and caused unstable boiler operations.
3. The stack had difficulty leak checking initially. The umbilical cord was found to be leaking at the cold box connection.
4. The west inlet hot box hit the ladder cage and caused it to be damaged. This resulted in the west inlet having to change out all glass ware and starting over. This caused us to be about an hour behind the other stations.
5. A unit went down at Plant X causing Harrington to pick up some load. The load was raised from 350 MW for a short while and then returned to 350 MW.

6. The east inlet sampled the last point of the first point for only two minutes. This was noted and was connected on the remaining ports.
7. Halfway through the first run on the west inlet the fitted disc collapsed causing the filter to break and allow particulate to enter the umbilical cord and impingers. As a result the test run was aborted at this sample station.

Representative Field Notes

BAGHOUSE STUDY
PHASE IV

1. SO₂ on Tuesday OK; particulate on Wednesday Run #1 west inlet at end of first port, we had to replace filters because of a wrinkle that developed. No particulate was expected to have passed filter; Run #2 - prior to sampling the west inlet, we tore the filter and interchanged reagents in impingers. Both hot and cold boxes were replaced prior to sampling. The east inlet filter became torn during the second part of the second point. The filters assembly was replaced at that point and the run resumed. Only a small amount of dust leaked past the filter; Run #3 - No significant problems.
2. See Richard Chambers report for baghouse status.
3. Prior to the testing on Tuesday, five bags in compartment 27 were found to have holes. This caused the opacity to be elevated. During the testing on TW&T the opacity was running about 8% and an occasional 10%.